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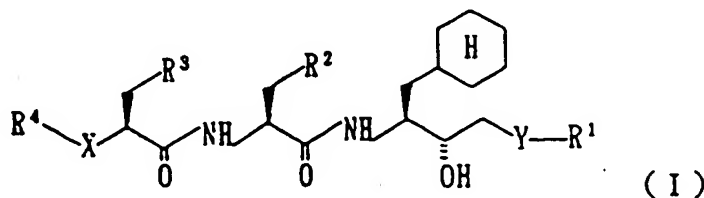
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(54) Renin inhibiting dipeptide derivatives, their preparation and pharmaceutical preparations containing them.

(57) A novel dipeptide derivative of the following formula (I), which compound is capable of inhibiting the enzymatic activity of renin and thereby depressing the renin-angiotensin system and lowering the blood pressure, is provided.



wherein:

R¹ is C₁-C₁₂ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₁₀ cycloalkyl, aryl, or heterocyclic radical;
R² is carbamoyl, aryl, 5- or 6-membered heterocyclic radical, C₁-C₁₂ alkyl-S-, C₁-C₁₂ alkyl-S-CH₂-, or C₃-C₁₀ cycloalkyl-S-;

R³ is aryl or 5- or 6-membered heterocyclic radical;

R⁴ is R⁴-SO₂ or R⁴-CO;

R⁴ is aryl, C₁-C₁₂ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl; C₃-C₁₀ cycloalkyl, or heterocyclic radical;

X is CH₂, NH, O, or S; and

Y is CO or NHSO₂, wherein R¹, R², R³ and R⁴ each may be substituted with one to three substituents selected independently from a group consisting of hydroxy; halogen; trifluoromethyl; -CN; heterocyclic radical; C₁-C₆ alkyl; C₃-C₁₀ cycloalkyl; -O-C₁-C₆ alkyl; C₁-C₆ alkylenedioxy; -CO-O-C₁-C₆ alkyl; -NHCO-C₁-C₆ alkyl; -S-C₁-C₆ alkyl; -SO-C₁-C₆ alkyl; -SO₂-C₁-C₆ alkyl; -NHSO₂-C₁-C₆ alkyl; -NR⁵R⁶; -O-CO-NR⁵R⁶; -CO-NR⁵R⁶; -O-C₁-C₆ alkyl NR⁵R⁶; R⁵ and R⁶ are independently hydrogen,

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formyl or C₁-C₈ alkyl, or R⁴ and R⁶, when taken together with the nitrogen to which they are attached, form a cyclic amino group, or an acid addition salt thereof.

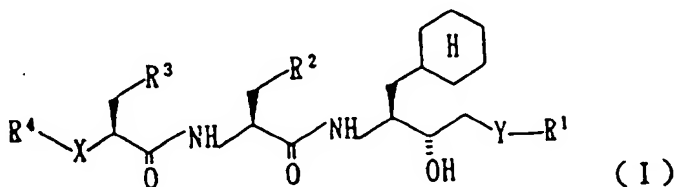
This invention relates to dipeptide derivatives capable of inhibiting renin activity, processes for their production and pharmaceutical preparations comprising them.

Renin (EC3.4.23.15) is a protease which catalyzes the hydrolysis of angiotensinogen into angiotensin I. The angiotensin I is a biologically inactive decapeptide, though it is enzymatically converted into angiotensin II by an angiotensin converting enzyme in pulmonal vascular endotheliocytes. This system is "the renin-angiotensin system". The angiotensin II induces hypertension through at least two routes, that is, contractive action on smooth muscles of peripheral vasculars and stimulation of secretion of adrenal hormone which inhibits sodium ion excretion. More particularly, it stimulates the secretion of aldosterone, an inhibitor of the excretion of Na⁺ ion, resulting in an increase of the volume of extracellular body fluid, rich is one of the causes of hypertension. Accordingly, compounds capable of depressing or inhibiting the renin-angiotensin system are expected to be potent anti-hypertensive substance. Many peptide analogues which seemed to be useful in the regulation of hypertensive diseases on the basis of renin-inhibiting activity have been developed and disclosed [for example, USP 4656269, EP-A-274259 and AU-A-8822959].

As mentioned above, the renin inhibitor inhibits the synthesis of Angiotensin I thereby depressing the renin-angiotensin system and lowering blood pressure. Owing to the physiological activity, renin inhibitors have been used in the treatment of hypertension. However, since hypertension is one of the most common disorders and causes many serious conditions and diseases, a development of more and more novel anti-hypertensive substances including renin inhibitors has been demanded to treat hypertension effectively.

The present inventors have now discovered a class of novel dipeptide compounds capable of inhibiting the catalytic activity of renin both *in vitro* and *in vivo*.

In particular, the present invention provides a dipeptide derivative of formula (I):



wherein:

R¹ is C₁-C₁₂ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₃-C₁₀ cycloalkyl, aryl, or heterocyclic radical;

R² is carbamoyl, aryl, 5- or 6-membered heterocyclic radical, C₁-C₁₂ alkyl-S-, C₁-C₁₂ alkyl-S-CH₂-, or C₃-C₁₀ cycloalkyl-S-;

R³ is aryl or 5- or 6-membered heterocyclic radical;

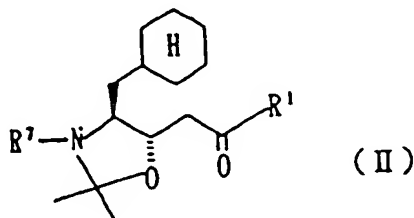
R⁴ is R^{4'}-SO₂ or R^{4'}-CO;

R^{4'} is aryl, C₁-C₁₂ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₃-C₁₀ cycloalkyl, or heterocyclic radical;

X is CH₂, NH, O, or S; and

Y is CO or NHSO₂, wherein R¹, R², R³ and R^{4'} each may be substituted with one to three substituents selected independently from a group consisting of hydroxy; halogen; trifluoromethyl; -CN; heterocyclic radical; C₁-C₈ alkyl; C₃-C₁₀ cycloalkyl; -O-C₁-C₈ alkyl; -S-C₁-C₈ alkyl; -SO-C₁-C₈ alkyl; -SO₂-C₁-C₈ alkyl; C₁-C₈ alkylenedioxy; -CO-O-C₁-C₈ alkyl; -NHCO-C₁-C₈ alkyl; -NHSO₂-C₁-C₈ alkyl; -NR⁵R⁶; -O-CO-NR⁵R⁶; -CO-NR⁵R⁶; -O-C₁-C₈ alkyl NR⁵R⁶; R⁵ and R⁶ are independently hydrogen, formyl or C₁-C₈ alkyl, or R⁵ and R⁶, when taken together with the nitrogen to which they are attached, form a cyclic amino group or an acid addition salt thereof.

In another aspect the present invention also provides a compound of formula (II):



wherein, R¹ is as defined above, and R⁷ is hydrogen or an amino protecting group, which compound is useful as an intermediate for the production of the compound of formula (I).

For the purpose of the present invention, as disclosed and claimed herein, the following terms are defined as below.

The term " C_1 - C_{12} alkyl" refers to a straight or branched saturated hydrocarbon radical having one to twelve carbon atoms, including methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *s*-butyl, *t*-butyl, *n*-pentyl, isopentyl, 2-methylbutyl, *t*-pentyl, neopentyl, isopentyl, 1-ethylpropyl, *n*-hexyl, isohexyl, heptyl, octyl, nonyl, decyl, undecyl, dodecyl, and the like.

The term " C_1 - C_6 alkyl" refers to a straight or branched saturated hydrocarbon radical having one to six carbon atoms as defined above.

The term " C_2 - C_6 alkenyl" refers to a straight or branched unsaturated hydrocarbon radical having two to six carbon atoms and one or more double bonds, including vinyl, allyl, 1-propenyl, isopropenyl, 2-butenyl, 1,3-butadienyl, 2-pentenyl, 1-hexenyl, and the like.

The term " C_2 - C_6 alkynyl" refers to a straight or branched unsaturated hydrocarbon radical having two to six carbon atoms and one or more triple bonds, including ethynyl, 1-propynyl, 2-propynyl, 2-butyne, 1,3-butadiynyl, 2-pentyne, 1-hexynyl, and the like.

The term " C_1 - C_6 alkylenedioxy" refers to methylenedioxy, ethylenedioxy, triethylenedioxy, tetramethylenedioxy, pentamethylenedioxy, hexamethylenedioxy, and the like.

The term " C_3 - C_{10} cycloalkyl" refers to cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, cyclononyl, cyclodecyl, and the like.

The term "aryl" refers to aryl radicals having 6 to 10 carbon atoms, including phenyl, indenyl, naphthyl, and the like.

The term "halogen" refers to halogen atoms such as fluorine, chlorine, bromine, and iodine.

The term "cyclic amino" refers to monocyclic or bicyclic amino groups such as pyrrolidino, 2-pyrrolidinyl, piperidino, 1-piperazinyl, 1-indolyl, 2-indolyl, morpholino, and the like.

The term "heterocyclic group" refers to a group of saturated or unsaturated monocyclic or condensed ring which contains one or more heteroatoms selected from nitrogen, oxygen and sulfur. Examples of heterocyclic groups include, for example, 2-thienyl, 3-thienyl, 2-furyl, 3-furyl, 2-pyrrolyl, 3-pyrrolyl, 2-imidazolyl, 1-pyrazolyl, 2-pyridyl, 3-pyridyl, 4-pyridyl, 2-pyrimidyl, 3-pyridazinyl, 2-pyrazinyl, 3-triazolyl, 2-thiazolyl, 4-thiazolyl, 5-tetrazolyl, 3-isothiazolyl, 2-pyrrolidinyl, 2-imidazolidinyl, 4-pyrazolidinyl, 4-piperidyl, 2-piperidinyl, 4-indolyl, 7-indolyl, 5-quinolyl, 8-quinolyl, 8-isoquinolyl, and the like.

The term "5- or 6-membered heterocyclic groups" refers to 5- or 6-membered heterocyclic groups as defined above.

The term "carbamoyl" refers to carbamoyl or carbamoyl substituted with one or two substituents selected from a group consisting of C_1 - C_6 alkyl or C_3 - C_{10} cycloalkyl, for example, carbamoyl, methylcarbamoyl, dimethylcarbamoyl, cyclohexylcarbamoyl, and the like.

In the definition of R^1 , preferred " C_1 - C_{12} alkyl" is methyl, ethyl, propyl, isopropyl, *t*-butyl, pentyl, hexyl, heptyl, or the like; preferred " C_1 - C_6 alkyl" is methyl, ethyl, propyl, isopropyl, *t*-butyl, or the like; preferred " C_2 - C_6 alkenyl" is vinyl, or the like; preferred " C_2 - C_6 alkynyl" is ethynyl, or the like; preferred " C_3 - C_{10} cycloalkyl" is cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, or the like; preferred "aryl" is phenyl, naphthyl, or the like. Preferred "heterocyclic group" is 5- or 6- membered heterocyclic group such as 2-thienyl, 2-furyl, 2-pyrrolyl, 2-thiazolyl, 4-thiazolyl, 5-tetrazolyl, 4-pyridyl, 5-pyrimidinyl, 2-pyrazinyl, 2-pyrrolidinyl, 4-piperidyl, or the like or condensed heterocyclic group such as 8-quinolyl, or the like.

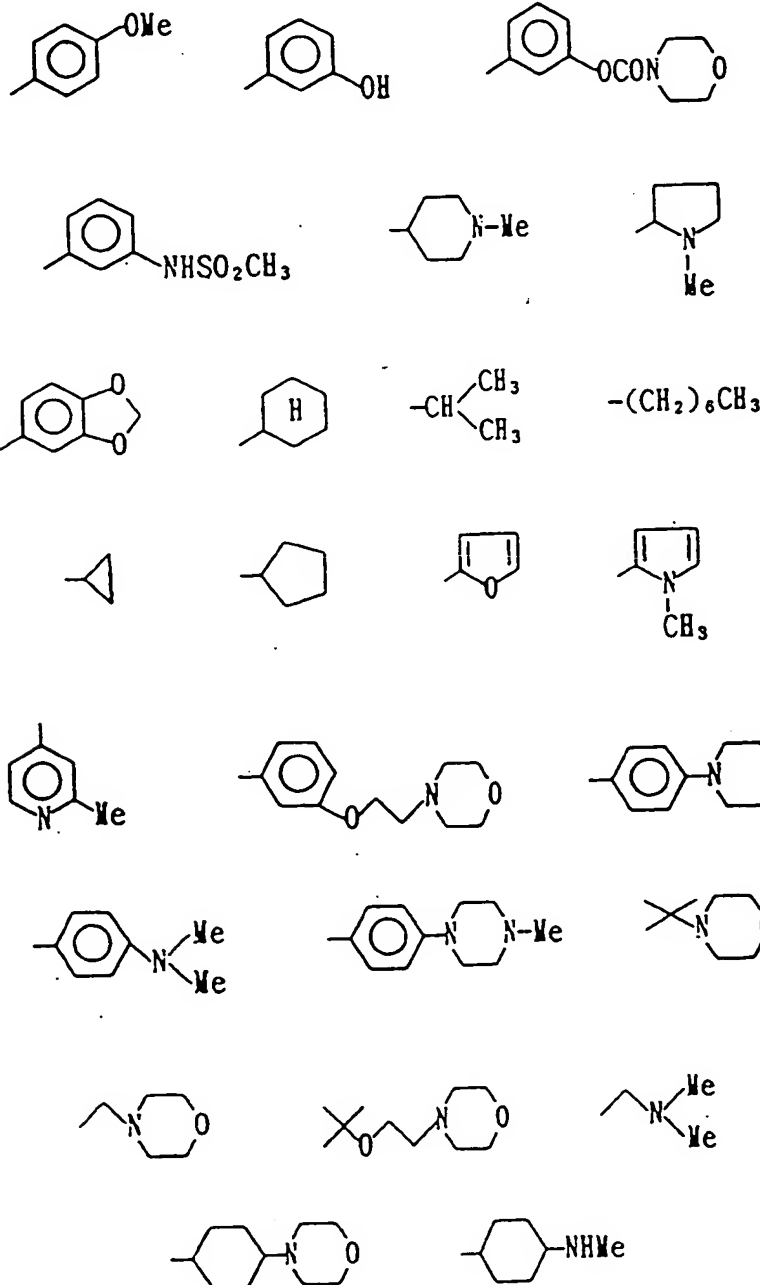
Examples of preferable R^1 include phenyl, *o*-tolyl, *p*-tolyl, *m*-tolyl, 2-chlorophenyl, 3-chlorophenyl, 4-chlorophenyl, 2,4-dichlorophenyl, 2,6-dichlorophenyl, 2-bromophenyl, 3-bromophenyl, 4-bromophenyl, 2,4-dibromophenyl, 2,6-dibromophenyl, 2-fluorophenyl, 3-fluorophenyl, 4-fluorophenyl, 2,4-difluorophenyl, 2,6-difluorophenyl, 2-tolylfluoromethyl, 3-tolylfluoromethyl, 4-tolylfluoromethyl, 2-hydroxyphenyl, 3-hydroxyphenyl, 4-hydroxyphenyl, 2-methoxyphenyl, 3-methoxyphenyl, 4-methoxyphenyl, 3,4-dimethoxyphenyl, 3,4-methylenedioxyphenyl, 3-methylaminophenyl, 3-(*N*-formyl)methylaminophenyl, 2-dimethylaminophenyl, 3-dimethylaminophenyl, 4-dimethylaminophenyl, 2-morpholinophenyl, 3-morpholinophenyl, 4-morpholinophenyl, 2-(4-methylpiperazino)phenyl, 3-(4-methylpiperazino)phenyl, 4-(4-methylpiperazino)phenyl, 2-acetamidophenyl, 3-acetamidophenyl, 4-acetamidophenyl, 2-methylsulfonylaminophenyl, 3-methylsulfonylaminophenyl, 4-methylsulfonylaminophenyl, 2-isopropoxycarbonylphenyl, 3-isopropoxycarbonylphenyl, 4-isopropoxycarbonylphenyl, 2-morpholinocarbonylphenyl, 3-morpholinocarbonylphenyl, 4-morpholinocarbonylphenyl, 2-morpholinocarbonyloxyphenyl, 3-morpholinocarbonyloxyphenyl, 4-morpholinocarbonyloxyphenyl, 2-morpholinoethoxyphenyl, 3-morpholinoethoxyphenyl, 4-morpholinoethoxyphenyl, 2-cyanophenyl, 3-cyanophenyl, 4-cyanophenyl, naphthyl, 2-pyrrolyl, 3-pyrrolyl, 1-methyl-2-pyrrolyl, 5-tetrazolyl, 1-methyl-5-tetrazolyl, 2-furyl, 3-furyl, 2-thienyl, 3-thienyl, 2-pyridyl, 3-pyridyl, 4-pyridyl, 1-methyl-4-pyridyl, 2-methyl-4-pyridyl, 3-methyl-4-pyridyl, 1-chloro-4-pyridyl, 2-chloro-4-pyridyl, 3-chloro-4-pyridyl, 1-fluoro-4-pyridyl, 2-fluoro-4-pyridyl, 3-fluoro-4-pyridyl, 2-pyrimidinyl, 5-pyrimidinyl, 2-pyrazinyl, 2-pyrrolidinyl, 3-pyrrolidinyl, 1-methyl-2-pyrrolidinyl, 1-methyl-3-

pyrrolidinyl, 2-piperidyl, 3-piperidyl, 4-piperidyl, 1-methyl-2-piperidyl, 1-methyl-3-piperidyl, 1-methyl-4-piperidyl, 8-quinolyl, methyl, ethyl, isopropyl, butyl, isobutyl, tert-butyl, pentyl, hexyl, heptyl, octyl, dimethylaminomethyl, morpholinomethyl, 1-morpholinoisopropyl, 1-morpholinoethoxyisopropyl, 1-piperidinomethyl, cyclopropyl, cyclopentene, cyclohexyl, cycloheptyl, cyclooctyl, 2-morpholinocyclohexyl, 3-morpholinocyclohexyl, 4-morpholinocyclohexyl, 2-methylaminocyclohexyl, 3-methylaminocyclohexyl, 4-methylaminocyclohexyl, 2-dimethylaminocyclohexyl, 3-dimethylaminocyclohexyl, 4-dimethylaminocyclohexyl, and the like.

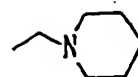
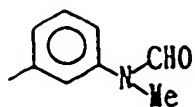
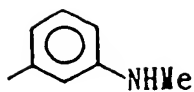
Examples of preferable R² include 5-membered heterocyclic groups containing two heteroatoms such as two nitrogen atoms, nitrogen and oxygen atoms, or nitrogen and sulfur atoms, for example, 4-imidazolyl, 4-thiazolyl, 4-oxazolyl, or the like, wherein said heterocyclic group may be substituted with methyl, ethyl, isopropyl, tert-butyl, amine, methylamine, dimethylamine, diethylamine, 1-pyrrolidinyl, piperidino, or the like; C1 - C12 alkyl-S- such as methylthio, ethylthio, cyclohexylthio, or the like; C1-C12 alkyl-S-CH₂- such as methylthiomethyl, or the like; carbamoyl or substituted carbamoyl such as methylcarbamoyl, dimethylcarbamoyl, or the like.

Examples of preferable R⁴ include sulfonyl or carbonyl substituted with methyl, ethyl, isopropyl, dimethylamino, tert-butyl, N-morpholino or N-morpholinomethyl, or the like.

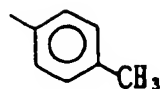
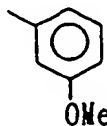
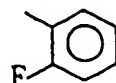
Examples of more preferable R¹ are shown below.



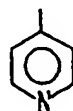
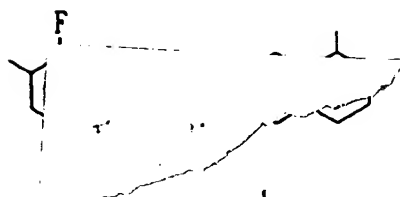
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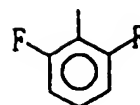
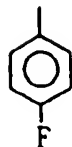
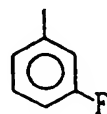
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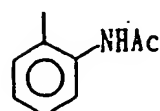
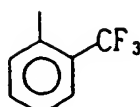
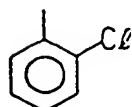
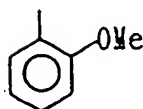
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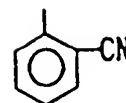
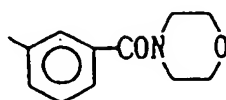
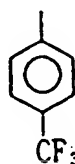
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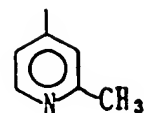
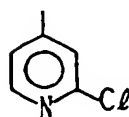
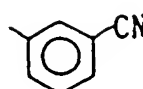
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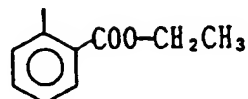
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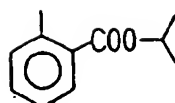
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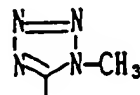
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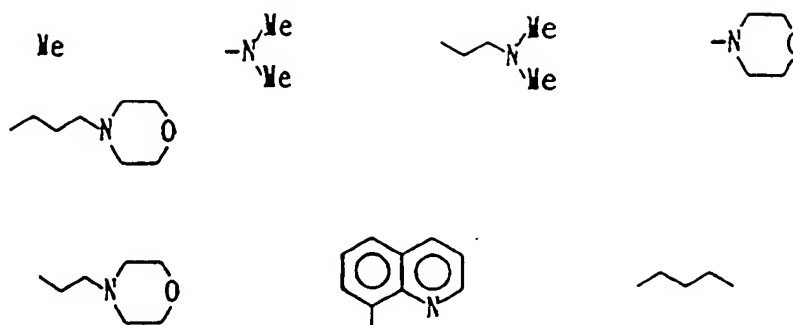
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Especially preferred compounds are those wherein R^2 is an optionally substituted 5- or 6-membered heterocyclic group; R^3 is an optionally substituted aryl; R^4 is morpholinosulfonyl; and X is NH .

The pharmaceutically acceptable acid addition salts of compounds of formula (I) include salts derived from a mineral acid such as hydrochloric acid, sulfuric acid, p-toluenesulfonic acids, or the like; carboxylic acid such as oxalic acid, maleic acid, citric acid, or the like. Preferable acid addition salts are those derived from mineral acid such as hydrochloric acid, sulfuric acid, toluenesulfonic acid, and the like.

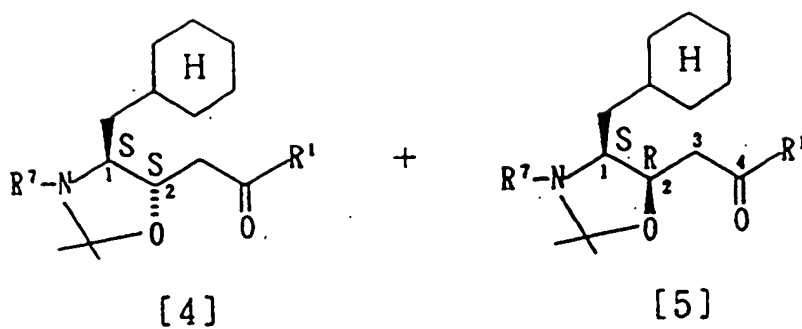
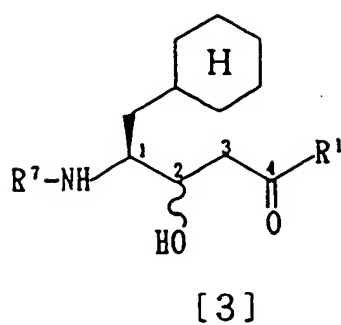
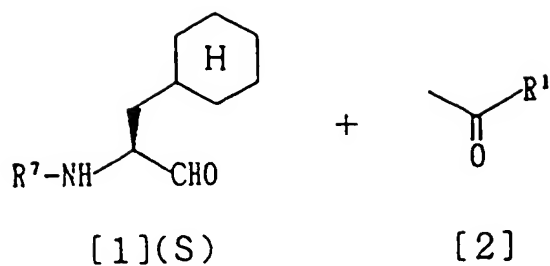
All the compounds of the present invention are novel and can be prepared according to either of two processes described below on the basis of what Y represents.

Process I

Preparation of compounds wherein Y is CO

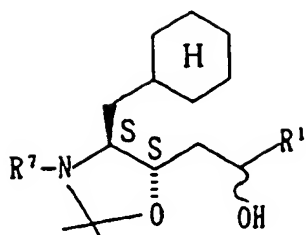
The process is schematically shown as below.

Step 1

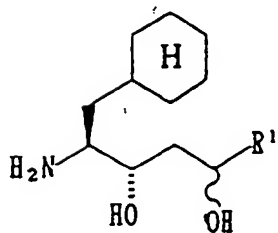


Step 2a

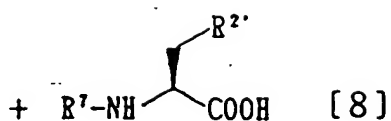
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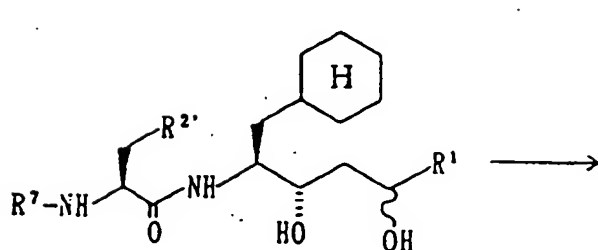
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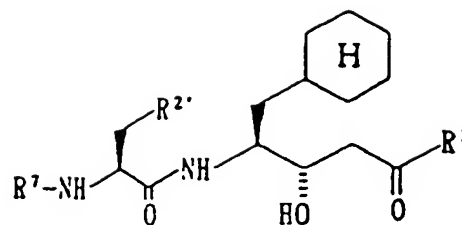
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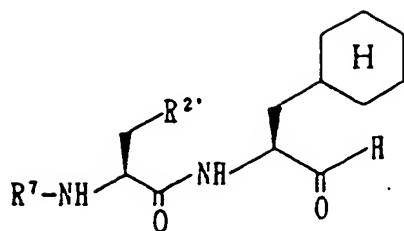
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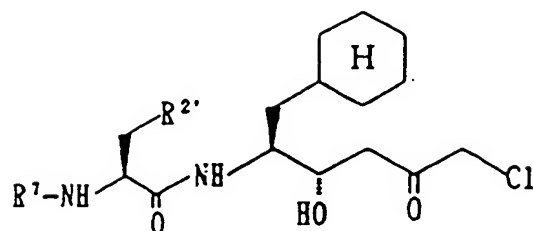
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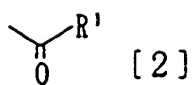
[10]

Step 2b

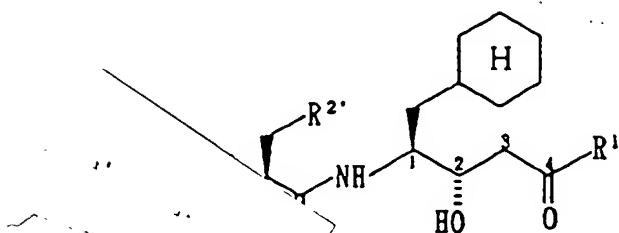
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Step 2c

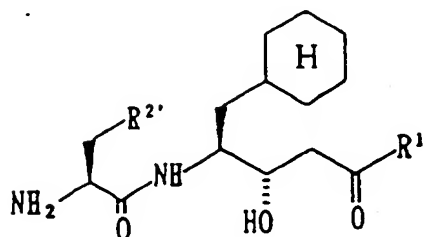
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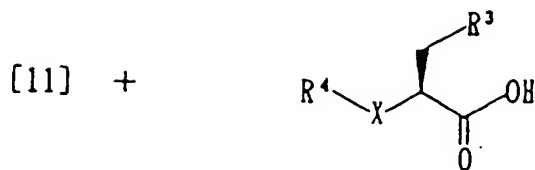
[2]



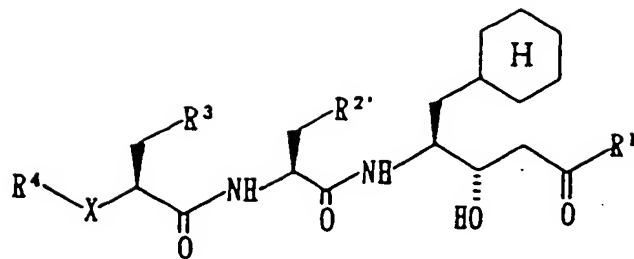
[10]

Step 3

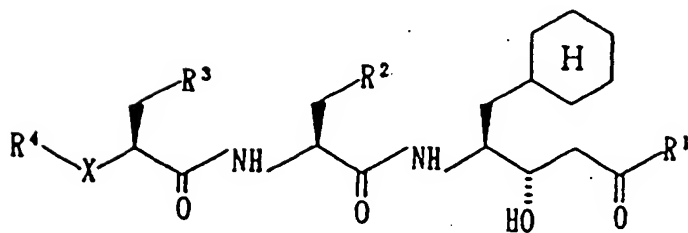
[11]

Step 4

[12] (S)



[13]



[1A]

In the above reaction schemes, R^1 , R^2 and R^3 are as defined above, $R^{2'}$ is optionally protected R^2 and R^7 is amino-protecting group.

The amino protecting group which is shown by R^7 can be selected from those groups generally used in peptide synthesis. Examples of amino protecting groups include benzyloxycarbonyl (it is referred to as Z), 2,6-dichlorobenzyloxycarbonyl ($Z(Cl)_2$), 4-nitrobenzyloxycarbonyl ($Z(NO_2)$), 4-methoxybenzyloxycarbonyl

(Z(OMe)), t-butoxycarbonyl (Boc), t-amylloxycarbonyl (Aoc), isobomylloxycarbonyl, adamantyloxycarbonyl (Adoc), 2-(4-biphenyl)-2-propyloxycarbonyl (Bpoc), 9-fluorenylmethoxycarbonyl (Fmoc), methylsulfonylethoxycarbonyl (Msc), trifluoroacetyl, phthalyl, formyl, 2-nitrophenylsulfonyl (NPS), diphenylphosphinothioyl (Ppt), dimethylphosphinothioyl (Mpt), and the like.

Examples of the optionally protected R^2 shown by R^2 are 4-imidazolyl, 4-aminothiazolyl and R^2 as defined above, which are optionally protected with a group selected from benzyl (Bzl), benzyloxycarbonyl (Z), toluenesulfonyl (tosyl or Ts), trimethylsilyl (trityl, Trt), dinitrophenyl (Dnp), 2,2,2-trifluoro-1-benzyloxycarbonylaminoethyl (Tfz), 2,2,2-trifluoro-1-t-butoxycarbonyl (TfBoc), adamantyloxycarbonyl (Adoc), piperidinocarbonyl, t-butoxycarbonyl(Boc), and the like.

Step 1

1. Preparation of Compound [3] by Aldol Reaction

a) The optically active aldehyde [1], a required starting compound, can be prepared from, for example, Boc-L-phenylalanine using any of known methods described in literatures such as ¹⁾ (T. Shioiri et al., J. Org. Chem. 52:1252 (1987) and J. Boger et al., J. Med. Chem. 28:1779 (1985)).

The aldol condensation between an aldehyde [1] and a ketone [2] is carried out by a novel stereoselective method of the present invention. The reaction is conducted using metal amide, as a base, in an organic solvent in the presence of a crown ether at a temperature of about -78°C. Amides which may be used include sodium bis-trimethylsilylamide ($\text{NaN}(\text{TMS})_2$), potassium bis-trimethylsilylamide ($\text{KN}(\text{TMS})_2$), lithium diisopropylamide, lithium bis-trimethylsilylamide, and the like. Crown ethers which may be used include 15-crown-5, 12-crown-4, 18-crown-6, and the like. Although all the combinations of amides and crown ethers described above are suited for the stereoselective aldol reaction of the invention, certain combinations are especially preferable in connection with the stereoselectivity of the product [3] which is expressed by the ratio of the product of 2S form to 2R form, i.e., diastereo-selectivity, 2S:2R. Thus, $\text{NaN}(\text{TMS})_2$, when used in association with 15-crown-5, gives the most favorable result shown by the 2S:2R value of about 2.4 to about 16.0, while other amides, when used alone or in combination with a crown ether, give inferior results shown by the 2S:2R value of less than 2.

Solvents which may be used include ethers such as diethyl ether, tetrahydrofuran (THF), dimethoxyethane, and the like with a preference for THF. When toluene is used, the stereoselectivity may be relatively decreased.

The reaction is carried out at a temperature ranging from about -20 to about -100°C, preferably about -78°C.

b) Alternatively, the stereoselective aldol condensation reaction can be carried out using metal alkoxide as a base in an inert solvent in the presence of a quarternary ammonium salt at a temperature of about -78°C.

Metal alkoxides which may be used include potassium t-butoxide (t-BuOK), potassium t-amyloxide ($\text{Et}(\text{Me})_2\text{COK}$) or sodium ethoxide (EtONa), and the like.

Quarternary ammonium salts which may be used include tetrabutyl ammonium bromide ($(\text{n-Bu})_4\text{NBr}$), tetramethyl ammonium bromide ($(\text{Me})_4\text{NBr}$), tributylbenzylammonium bromide ($\text{Bn}(\text{n-Bu})_3\text{NBr}$), and the like. All the reagents are suited to the stereoselective aldol reaction of the invention and the best result can be obtained by the combination of t-BuOK and $\text{n-Bu}_4\text{NBr}$ giving the 3S/3R value of about 3.3 - 6.5. This method is useful even in the absence of quarternary ammonium salt and gives the ratio of 3S/3R of about 3 to 5.

Solvents which may be used include THF, toluene, dichloroethane, dichloromethane, and the like with a preference for dichloromethane. When THF or toluene is used, the stereoselectivity may be decreased. The reaction can be conducted under a similar temperature as described in above a).

2) Separation of Stereoisomer (1S, 2S) [4]

The desired stereoisomer [3]-(2S) can be separated from a mixture of isomers shown by formula [3] by a known resolving procedure, for example, a column chromatography on silica gel. For the purpose of the invention, the desired isomer can be conveniently separated by reacting the mixture [3] with 2-methoxypropene or 2,2-dimethoxypropane in the presence of a catalytic amounts of p-toluene sulfonic acid or pyridinium p-toluene sulfonate in a solvent such as THF or dichloroethane at a temperature ranging from room temperature to the refluxing temperature for about 1 to 8 hours to obtain a product containing a mixture of ring-closed compounds [4] and [5] which differ in the crystallizing properties from a certain solvents. Thus, when the product is recrystallized from ethyl acetate or diisopropyl ether in which the desired stereoisomer [4] is hardly soluble and the undesired isomer [5]-(2R) is soluble, the former can be separated as a crystalline solid, while the latter remains in the mother liquor. A column chromatography on, for example, silica gel, can be used when the compound [4] is not easily separated by recrystallization. The so obtained compound [4] in (1S, 2S) form is a novel and useful compound as an intermediate for the production of the compound (I).

Alternatively, the product [3], without further treatment to form acetonide, can be directly subjected to a column chromatography on silica gel to yield the stereoisomer [3]-(2S), which is then converted into dihydric alcohol of formula [7].

5 Step 2a

Before the deprotection of C1 amino group, the compound [4] should be reduced to avoid the possibility of ring closing reaction between the deprotected amino group and the C4 carbonyl group. The reducing reaction can be carried out using any of known methods in the art. However, it is efficiently conducted by reacting a solution of the ketone [4] in ethanol, methanol, THF or toluene with a reducing reagent such as sodium borohydrate, L-selectride or Red-Al at room temperature or under cooling for about 0.5 to 2 hours. Preferably, the latter reagent is used slightly in excess, that is, about 1.0 to 1.3 mole to 1.0 mole of ketone [4]. The resultant product [6], a mixture of diastereoisomers (1:1 to 3:1), is used in the next deprotection step without further purification.

The deprotection of amino group can be carried out using any of following procedures. When the protecting group is Boc, and the like, the compound [6] is deprotected by dissolving into THF or dioxane, adding 6N HCl thereto, and stirring at room temperature for about 1 to 4 hours. Alternatively, the compound [6] is treated with an acid such as aluminium chloride, trifluoroacetic acid or formic acid in the presence of anisole to yield the dihydric aminoalcohol [7].

When the protecting group is a member of benzyloxycarbonyl groups such as benzyloxycarbonyl (hereinafter, it is referred to as Z), 2,6-dichlorobenzyloxycarbonyl (Z(Cl)₂), or 4-nitrobenzyloxycarbonyl ((Z(NO₂))), the deprotection can be effected by catalytic reduction using palladium-containing catalyst, and the like. When the protecting group is Fmoc (9-fluorenylmethoxycarbonyl), Msc (methylsulfonylethoxycarbonyl), or the like, the deprotection can be effected by treating the compound [6] by piperidine, diethylamine, or the like.

The resulting dihydric alcohol of formula [7] is subjected to the next condensation reaction without purification. The condensation can be carried out using any procedure generally used in the field of peptide synthesis. For example, to a solution of compound [7] in an appropriate solvent such as dichloromethane is added commercially available N-Boc-amino acid [8] or its DCHA salt, and the mixture is allowed to react at room temperature for about 1 to 4 hours in the presence of a slight excess of a coupling reagent such as 1.0 to 1.3 mole equivalent of diethyl cyanophosphosphate (DEPC) and, if desired, a tertiary amine such as N-methyl morpholine to obtain a coupled compound [9]. Examples of coupling reagents are DCC, EDC, DEPA, BOP, DCC-HOBT, DCC-HOSu, ethyl chlorocarbonate, isobutyl chlorocarbonate, isopropyl chlorocarbonate, diethyl chlorophosphate, diphenyl chlorophosphate, 2-chloro-4,6-dimethoxy-1,3,5-triazine, and the like. The compound [8] may be protected at the heterocyclic ring with a protecting group generally used in the field of peptide synthesis.

The resultant diastereoisomer [9] is also converted into the corresponding ketone [10] without separation by dissolving the compound [9] into dichloromethane or DMF, adding about 3 to 10 times amounts of active manganese dioxide to the mixture and reacting at room temperature for 2 to 8 hours. This reaction proceeds very smoothly when fine crystal starting material [9] is used. The characteristic of this reaction is that the hydroxyl group at the C4 position of benzyl compound can be selectively oxidized.

40 Step 2b

Compound [10] can be also prepared through an aldol reaction according to a procedure described in step 1 from a starting compound [2] and a dipeptide aldehyde of formula [14] obtainable from a corresponding dipeptide alcohol in the same manner as that used for the preparation of compound [1]. The reaction however proceeds without stereoselectivity and differs from that of step 1 in this regard. The product being a 1:1 mixture of compound [10] in 2S and 2R isomers, chromatographic procedure is required for the separation of desired [10]-(2S)-isomer. The characteristic of the method of step 2b is that it is applicable when the method of above step 2a is not effective because a compound resists the selective oxidization with manganese dioxide.

50 Step 2c

The compound [10] can be prepared by reacting a chloromethyl ketone of formula [19] with an amine. The characteristic of the method of step 2c is that it is useful in the introduction of N-substituted methylketone residue to the C-terminal moiety.

Step 3

The deprotection of ketone compound [10] can be carried out in the same manner as described in the preparation of amino dihydric alcohol [7] from compound [6]. For example, when the protecting group is Boc, it is carried out by adding excess aluminium chloride to an anisole solution of compound [10] and stirring the mixture for about 1 to 3 hours at a temperature ranging from ice-cooled temperature to room temperature. The deprotection can also be effected by treating the compound [10] with either of excess trifluoroacetic acid in anisole or 6N HCl in THF to yield the desired compound [11]. The resultant ketone [11] with carbonyl group at the C4 position is novel and important as an intermediate for preparing the compound of formula (I) of the present invention.

Step 4

The compound [11] is reacted with sulfonyl propionic acid derivatives, N-sulfamyl, N-carbamoyl, or N-acyl amino acid of formula [12] which can be prepared according to a known method such as described in a literature (J.L. Stanton et al., J. Med. Chem. 31:1839 (1988)) under a condition for the coupling reaction and then deprotected if necessary to give the desired compound (IA) as the final product.

The coupling reaction is preferably conducted using 1.0 to 1.3 mole equivalent of diethyl cyanophosphonate (DEPC) in the presence of N-methyl morpholine (NMM) in a solvent such as dichloromethane at room temperature for about 1 to 8 hours. Examples of coupling reagents are DCC, EDC, DEPA, BOP, DCC-HOBt, ethyl chlorocarbonate, isobutyl chlorocarbonate, isopropyl chlorocarbonate, diethyl chlorophosphate, diphenyl chlorophosphate, 2-chloro-4,6-dimethoxy-1,3,5-triazine, and the like.

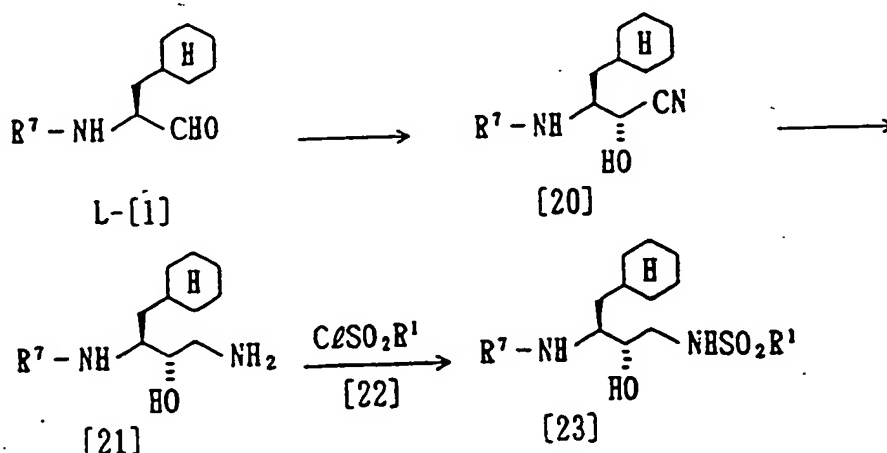
The deprotection of the compound [13] is carried out using any of known procedures depending on the protecting group. When the protecting group of R² is tosyl, it can be carried out by stirring a mixture of a solution of compound [13] in DMF in the presence of 5 to 12 mole equivalent of pyridinium hydrochloride at room temperature for about 1 to 4 hours. The deprotection can be effected by means of trifluoroacetic acid (at 15°C for about 30 minutes), HBr/acetic acid (at room temperature for about 30 minutes), conc. ammonia (at room temperature for about 1 hour), conc. HCl, or the like.

Process II

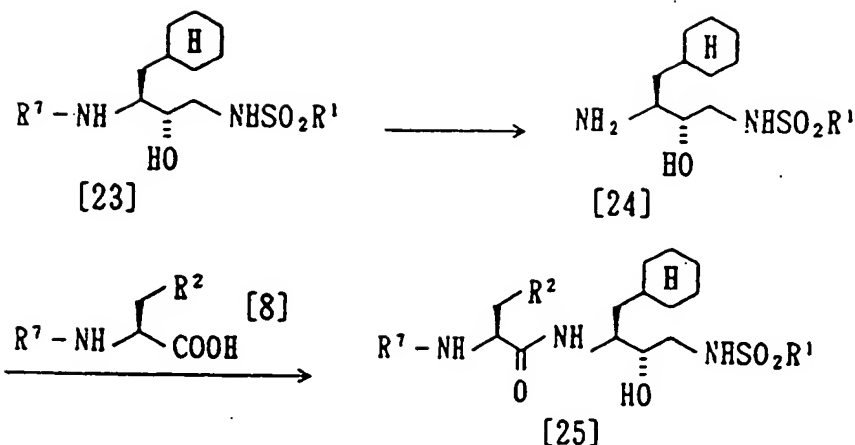
Preparation of compounds (I) wherein Y is NHSO₂

The process is schematically shown as below.

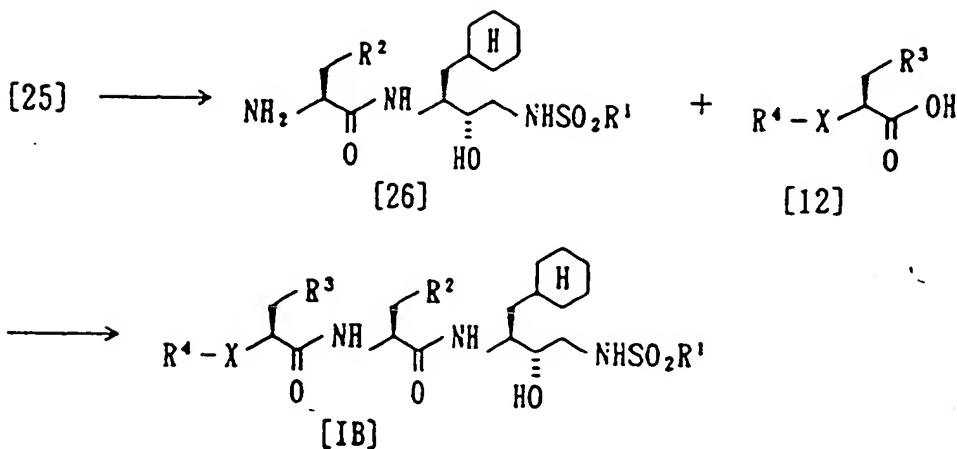
Step 1



Step 2



Step 3



In the above reaction schemes, R¹, R², R³, R⁴ and R⁷ are as defined above.

Step 1

The optically active aldehyde [1], a required starting compound, can be prepared in the same manner as described in above Process I.

The preparation of cyanhydrin [20] from aldehyde [1] is carried out substantially in accordance with a procedure described in the literature. Thus, the aldehyde [1] is allowed to react with an acidic sodium sulfite to obtain an additive, which is then reacted with KCN in ethyl acetate at room temperature to yield the cyanhydrin [20] stereoselectively (2R/2S = 3/1). The product is then resolved into each stereoisomer by column chromatography on silica gel. The desired (2R)-isomer is a crystalline solid and can be purified by recrystallization while the undesired (2R)-isomer is an oil. Therefore, alternatively, the desired product [20]-2R can be obtained conveniently by adding a seed crystal to the reaction mixture, collecting the precipitate, and recrystallizing from a solvent before subjecting to the chromatography.

The cyanhydrin [20] is then converted into an amino alcohol [21] by reducing the nitrile group. The reduction is carried out effectively by dissolving cyanhydrin [20] into an ethereal solvent, preferably THF, adding about 2 to 2.5 mole of lithium aluminium hydride thereto. The resulting amino alcohol [21] is then, without purification, reacted with sulfonyl chloride [22] to obtain sulfonyl amide of formula [23]. The reaction is conducted by reacting the amino alcohol [21] and sulfonyl chloride [22] in an appropriate solvent such as dichloromethane in the pre-

sence of tertiary amine such as triethylamine at room temperature for overnight.

Step 2

- 5 The deprotection of compound [23] can be carried out in a similar manner as described in the above Process I. The deprotected compound [24] is, without purification, dissolved into an appropriate solvent such as CH_3CN , or the like, and subjected to a condensation with N-protected-amino acid [8] in the same manner as the coupling reaction described in the above process I to yield a dipeptide analogue [25].

10 Step 3

- The compound [25] is then deprotected in the similar manner as that used for the deprotection of compound [23] in the above Process II, step 2. The product [26] is, without purification, subjected to the condensation reaction with a modified carboxylic acid [12] in exactly the same manner as described in Process I to obtain the final product [18].

- 15 As can be seen from the above reaction schemes, the present invention provides a dipeptide in which one peptide bond is formed through a coupling reaction between, for example, a free carboxyl group of an amino-protected amino acid and an amino group of an amino dihydric alcohol of formula [7] prepared from an oxazolidine derivative of formula [4]. The compound [4], an important intermediate for preparing the compound of formula (I), is obtained by a stereoselective aldol condensation method of the present invention. The other peptide bond is formed by a coupling reaction between a carboxylic group of, for example, sulfonyl propionic acid of formula [12] with a free amino group of a deprotected amino ketone [11] such as histidine as can be seen in the step 4.

- 25 As will be hereinafter described in the Experiment, the compounds of the invention have been demonstrated to be an effective renin inhibitor, whereby they suppress the renin-angiotensin system (one of the in vivo causes of hypertension) and lower blood pressure. The compounds of the invention are low in toxicity and useful in the treatment of hypertension or cardiac dysfunction through their renin inhibitory activity. The compounds may be administered either orally or parenterally. It is a characteristic benefit of the compounds that they are effective even when orally administered.

- 30 When the compounds of the invention are used to treat renin-associated disorders, a therapeutically effective amount of a compound of formula (I) is formulated into a composition of an appropriate form by known procedures using pharmaceutically acceptable carriers, diluents, or excipients. The administration may be conducted orally, intranasally, intravenously, subcutaneously, or the like.

- 35 For preparing composition for the administration, an active compound (I) is mixed with one or more standard adducts such as excipient, stabilizer, or inert diluent, and the like. The mixture is then formulated into an appropriate form such as a tablet, coated tablet, hard gelatin capsule, or an aqueous, alcoholic or oily suspension, or an aqueous, alcoholic or oily solution. Examples of inert excipients which can be used include various cyclodextrins, preferably β -cyclodextrin, acacia gum, magnesium carbonate, potassium phosphate, lactose, glucose, magnesium stearyl fumarate, starch, and the like. Either of dry or wet granules can be used. Examples of oily excipients or solvents include vegetable oil such as sunflower oil and fish liver oil.

- 40 For subcutaneous or intravenous administration, an active compound or a pharmaceutically acceptable salt thereof is dissolved, dispersed or emulsified into an appropriate solvent with the aid of any substances generally used in such a purpose, for example, solubilizing agent, emulsifying agent, or other adjuncts to obtain solutions, suspensions or emulsions.

- 45 Examples of appropriate solvents include water, physiological saline, alcohols such as ethanol, propanediol or glycerol, a sugar solution such as a solution of glucose or mannitol, or a mixture thereof, or Tween 80. Examples of solubilizing agents include above-mentioned cyclodextrins, preferably β -cyclodextrin.

The abbreviations used are as follows:

- 50 Boc = tertiary-butoxycarbonyl; Red-Al = sodium bis(2-methoxyethoxy)aluminium, L-Selectride = lithium tri-sec-butylborohydride; Boc His(Ts).DCHA = N^ε-Boc-N^α-tosyl-L-histidine dicyclohexylamine; BOP = benzotriazol-1-yl-oxy-tris-(dimethylamino)phosphoniumhexafluorophosphate;
DCC-HOBT = dicyclocarbodiimide-1-hydroxybenzotriazole;
DCC-HOSu = dicyclohexylcarbodiimide-N-hydroxysuccineimide;
DEPC = diethyl cyanophosphonate; NMM = N-methylmorpholine;
55 PPTS = pyridinium paratoluenesulphonate;
Tala = (4-thiazolyl)-L-alanine; rt = room temperature;
Ts = tosyl; TMS = trimethylsilane;
DMAP = 4-dimethylaminopyridine;

DCHA = Dicyclohexylamine;

DCC = Dicyclohexylcarbodiimide;

EDC = 1-Ethyl-3-(3dimethylaminopropyl)carbodiimide;

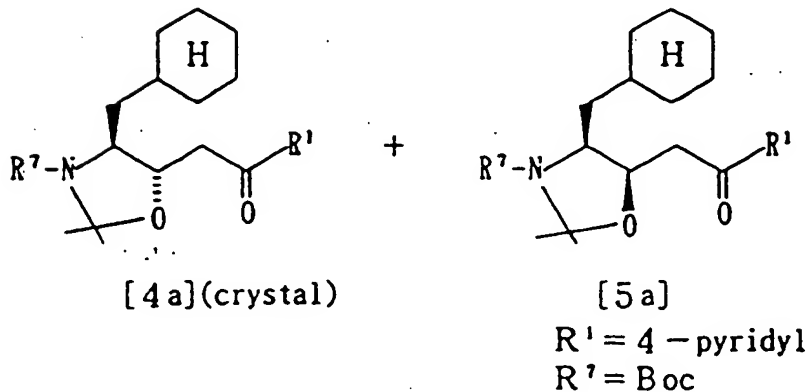
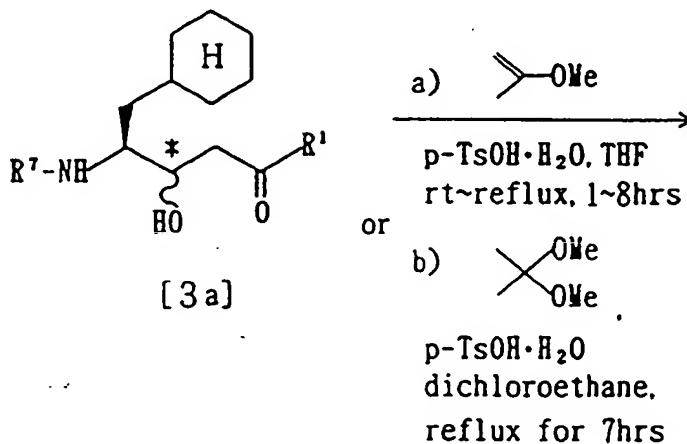
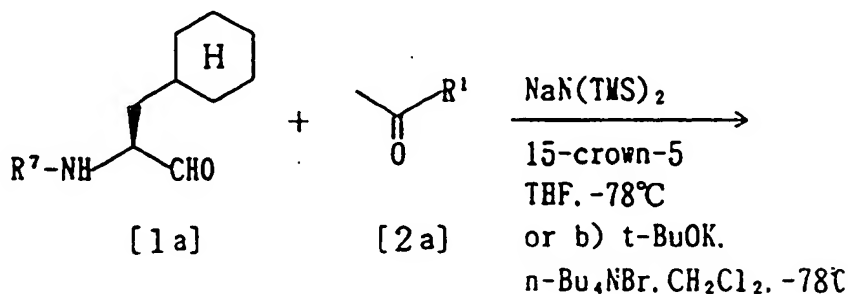
DEPA = Diethyl phosphorylazide;

5 BOP = Benzotriazol-1-yl-oxy-tris(dimethylamino)-phosphonium hexafluorophosphate

The following Examples further illustrate the compounds of the invention and the processes for preparing the same. The Examples are not intended to be limiting to the scope of the invention in any respect and should not be so construed. Unless otherwise noted, the NMR spectra were measured in CDCl_3 at 200 MHz (internal standard = TMS) and IR spectra in CHCl_3 . All amino acid used are L-isomers.

Preparation 1

3-Boc-4-(S)-cyclohexylmethyl-2,2-dimethyl-5(S)-[2-oxo-2-(4-pyridyl)ethyl]oxazolidine [4a]



1. a) To a 36ml (36mmol, 1.5eq) solution of 1N NaN(TMS)_2 in THF is added a solution of 4.34g (36mmol, 1.5 eq) of 4-acetylpyridine [2a] in 20ml of THF at -78°C over 10 minutes under nitrogen atmosphere. After 10 minutes stirring, a solution of 7.898g (36mmol, 1.5 eq) of 15-crown-5 in 10 ml of THF is added thereto and stirred

for 5 minutes. To the mixture is added 6.108g (24 mmol) of N-Boc-L-cyclohexylalaninal [1a] in 50ml of THF over 15 minutes and stirred for 1 hour at -78°C. The reaction mixture is added to a mixture of saturated aqueous solution of ammonium chloride and ethyl acetate with stirring and extracted three times with ethyl acetate. The extract is washed with saturated brine, dried over magnesium sulfate and concentrated to dryness *in vacuo*.
 5 The residue, upon purification by column chromatography on silica gel (eluent; dichloromethane/methanol = 98.5:1.5) gives N-Boc-1 (S)-cyclohexylmethyl-2-hydroxy-4-oxo-4-(4-pyridyl)butylamine [3a] (5.94g; yield = 66.0%) as a colorless powder. The product is a mixture of compound of 2(S)-isomer (desired isomer) and 2(R)-isomer (the ratio of 2(S) : 2(R) = 5.24 : 1).

b) To a stirring solution of 32g (125.3mmol) of N-Boc-L-cyclohexylalaninal [1a], 22.8g (188mmol, 1.5eq) of N-acetylpyridine, and 60.6g (188mmol, 1.5eq) of tetrabutyl ammonium bromide in 700ml of dichloromethane is added each one fourth portions of t-BuOK (21.1g in total, 188mmol, 1.5eq.) at 10 minutes interval under cooling at -78°C and the stirring is continued for another 1.5 hours at the same temperature. The reaction mixture is added to a mixture of saturated aqueous ammonium chloride and dichloromethane with stirring and extracted three times with dichloromethane. The extract is treated with citric acid to purify the basic substances to obtain
 15 a crude product [3a] (37g; yield = 79%; 2(S)/2(R) = 7 : 1).

2. a) To a solution of 5.908g (15.7mmol) of purified alcohol [3a] in 50ml of THF are added 2ml (20.9mmol, 1.3 eq) of 2-methoxypropene and 299mg (1.57mmol, 0.1eq) of p-toluenesulfonic acid monohydrate and the mixture is heated to reflux for 4 hrs. The reaction mixture is concentrated under reduced pressure, and the residue is alkalified with 4% sodium bicarbonate and extracted 3 times with dichloromethane. The extract is washed
 20 once with saturated brine, dried over magnesium sulfate, and concentrated to dryness. The residue is decolorized by column chromatography on silica gel using a short column (eluent; dichloromethane/acetonitrile = 5:1) and recrystallized from ethyl acetate to obtain 4.66g (yield = 68.6%) of the title compound [4a] as a colorless solid.

b) A mixture of 72g (195.6mmol) of the crude alcohol [3a], 150ml (122.0mmol, 6.2eq) of 2,2-dimethoxypropane and 2.73g (14.4mmol, 0.073eq) of p-toluenesulfonic acid monohydrate in 150ml of dichloroethane is heated to reflux for 16 hours. After cooling, the mixture is made basic with 4% aqueous sodium bicarbonate and extracted 3 times with dichloromethane. The extract is washed once with saturated brine, dried over magnesium sulfate, and concentrated to dryness *in vacuo*. The crude product, upon recrystallization from isopropyl
 25 ether, gives 23.5g (29.5%) of the compound [4a] as a white crystal. The mother liquor, when treated by column chromatography on 300g of silica gel (eluent; dichloromethane/ethyl acetate = 7:1) and recrystallized in the same manner as above, gives 2.5g (3.1%) of compound [4a].
 m.p. = 115 - 116°C

$[\alpha]_D^{25} = -18.5^\circ$ (C=1.0 CHCl₃; 23.5°C)

IR_{max}(CHCl₃): 1692, 1596, 1557, 1477, 1450, 1172, 1086 cm⁻¹

35 NMR δ (CDCl₃): 1.48(9H,s), 1.52(3H,s), 1.60(3H,s), 0.78-1.90(13H,m), 3.14(1H,dd,J=16.8,6.8Hz), 3.41(1H,dd,J=16.7,6.1Hz), 3.84(1H,m), 4.52(1H,t like m), 7.73(2H,m), 8.83(2H,m)

Elemental analysis (as C₂₄H₃₈N₂O₄)

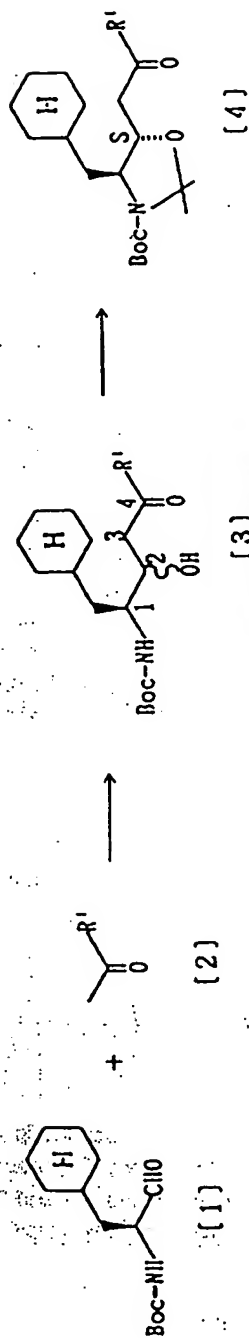
Calcd.(%): C:69.20; H:8.71; N:6.73

Found (%): C:69.20; H:8.75; N:6.76

40 Preparation 2 - 20

Compounds [4], the desired stereoisomers, were prepared according to the method described in above Preparation 1 by preparing compound [3a] and separating the desired isomer [3]-(S) therefrom. The results are
 45 shown in the following Table 1. Among the compounds listed in the Table 1, compound Nos. 13 and 14 are separated chromatographically because the corresponding compounds of formula [4] do not crystallize under the conditions used.

Table I



Compd. of Prep. No.	R'	[3]		mp $^{\circ}\text{C}$	[α] $_D^{25}$ C=1.0, CHCl ₃ ($^{\circ}\text{C}$)	Elemental analysis	Calcd.	Found	IR ν_{max}
		Yield%	C-2 S/R						
2	phenyl	71	3.1	111~ 113	-17.4 (23.5)	C ₂₁ H ₁₉ NO ₄	C: 72.25 H: 8.98 N: 3.37	C: 72.25 H: 8.99 N: 3.36	1686, 1650, 1582, 1478, 1450, 1172, 1088
3	o-fluorophenyl	68	4.8	95~ 97	-18.5 (24.0)	C ₂₈ H ₂₃ NO ₄ F	C: 69.25 H: 8.37 N: 3.23 F: 4.38	C: 69.12 H: 8.10 N: 3.23 F: 4.45	1686, 1610, 1577, 1480, 1453, 1173, 1100, 1086, 990, 848
4	m-methoxyphenyl	75	2.7	117~ 119	-6.2 (23.5)	C ₂₇ H ₂₅ NO ₄	C: 70.08 H: 8.82 N: 3.14	C: 70.05 H: 8.74 N: 3.15	1689, 1600, 1585, 1488, 1465, 1456, 1430, 1394, 1369, 1290, 1255, 1172, 1139, 1088, 1050
5	p-methoxyphenyl	78	2.4	132~ 134	-23.5 (24.0)	C ₂₈ H ₂₇ NO ₄	C: 72.69 H: 9.15 N: 3.26	C: 72.66 H: 9.08 N: 3.20	1687, 1610, 1573, 1480, 1450, 1174, 1088,
6	2,4-difluorophenyl	91	4.8	135~ 137	-19.1 (23.5)	C ₂₈ H ₂₃ NO ₄ F ₂	C: 66.49 H: 7.81 N: 3.10 F: 8.42	C: 66.31 H: 7.82 N: 3.04 F: 8.38	1687, 1612(1595), 1498, 1477, 1450, 1430, 1172, 1140, 1098, 971, 855

Table 1 (continued)

Compd. of Prep. No.	R'	[3]		mp°C	[α] _D ²⁰ C=1.0, CHCl ₃ (°C)	Elemental analysis	Calcd.	Found	IR ν cm ⁻¹ max
		Yield%	C-2 S/R						
7	1-naphthyl	90	1.7	127~ 128	-11.7 (24.0)	C ₂₁ H ₁₃ NO ₄	C: 74.81 H: 8.44 N: 3.01	C: 74.84 H: 8.43 N: 3.06	1687, 1595, 1508, 1477, 1449, 1393, 1379, 1368, 1250, 1172, 1138, 1098, 1085
8	3-thienyl	80	2.7	113~ 114	-13.5 (25)	C ₂₂ H ₁₃ NO ₄ S	C: 65.52 H: 8.37 N: 3.32 S: 7.61	C: 65.75 H: 8.28 N: 3.31 S: 7.57	1685, 1510, 1477, 1450, 1172, 1088
9	2-thiazolyl	75	16	128~ 129	-10.7 (23.5)	C ₂₂ H ₁₃ N ₂ O ₄ S	C: 62.53 H: 8.11 N: 6.63 S: 7.59	C: 62.28 H: 7.79 N: 6.53 S: 7.36	1690, 1480, 1448, 1170, 1075, 945

Table 1 (continued)

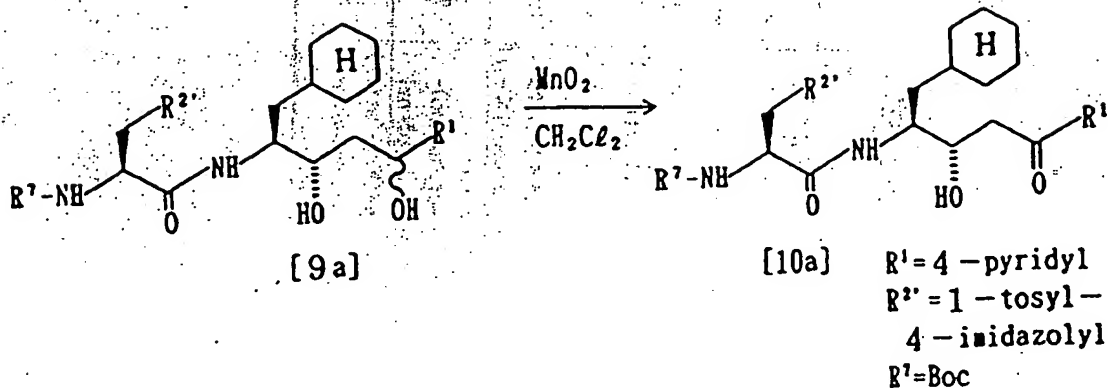
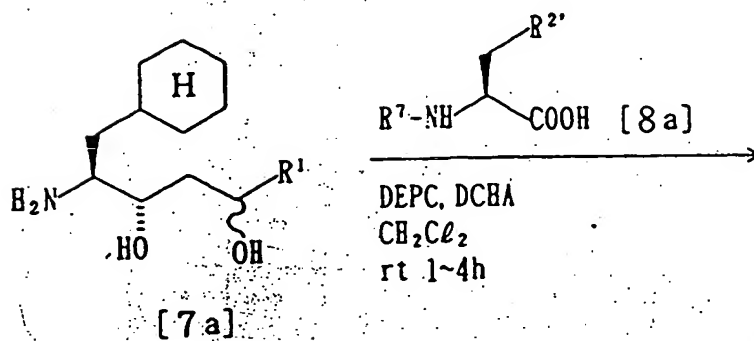
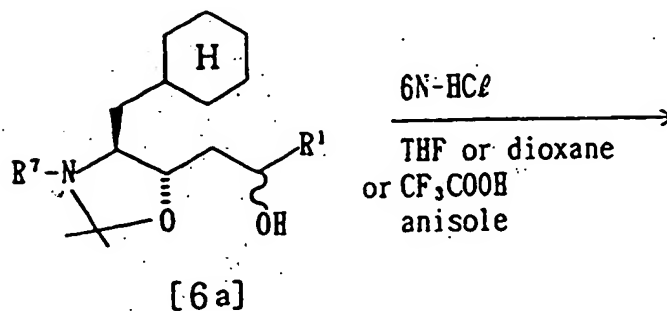
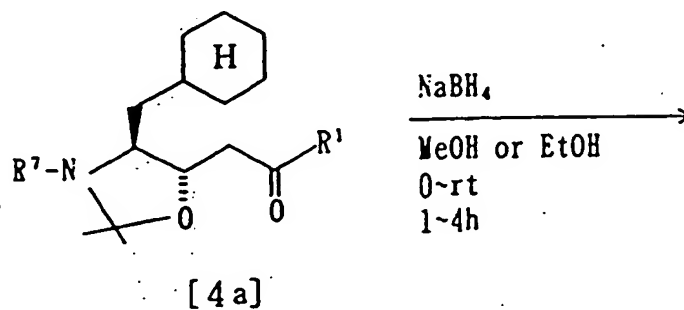
Compd. of Prep. No.	R'	[3]		Yield%	mp°C	[α] _D ²⁵ C=1.0, CHCl ₃	Elemental analysis	[3](S) or [4]		IR ν _{max} ^a or NMR(δ)
		Yield%	C-2 S/R					Calcd.	Found	
10	m-fluorophenyl	82	3.9	71	103~ 105	-15.7°	C ₂₁ H ₁₃ N ₁ O ₄ F	C: 69.25 H: 8.37 N: 3.23 F: 4.38	C: 69.36 H: 8.41 N: 3.25 F: 4.22	1690, 1610, 1590, 1485, 1475, 1443, 1392, 1170, 1086
11	p-fluorophenyl	83	2.8	61	137~ 138	-15.7°	C ₂₁ H ₁₃ N ₁ O ₄ F	C: 69.25 H: 8.37 N: 3.23 F: 4.38	C: 69.14 H: 8.35 N: 3.14 F: 4.41	1685, 1600, 1505, 1475, 1450, 1392, 1170, 1155, 1085
12	2,6-difluorophenyl	88	13.0	83	51~ 54	-18.8°	C ₂₁ H ₁₁ N ₁ O ₄ F ₂	C: 66.50 H: 7.81 N: 3.10 F: 8.41	C: 66.40 H: 7.79 N: 3.34 F: 8.69	1691, 1624, 1588, 1467, 1450, 1394, 1369, 1279, 1174, 1139, 1089, 1030, 982, 860
13	o-acethoxyphenyl	79	2.8	[3](S) 53						0.75-1.93(13H, m), 1.45(9H, s), 3.10(1H, dd, J=9.9, 18.3Hz), 3.70(1H, m), 4.16(1H, m), 4.82(1H, d, J=10Hz), 7.00(2H, m), 7.50(1H, td, J=2.5, 7Hz), 7.75(1H, dd, J=2.5, 7Hz)
14	o-chlorophenyl	86	3.0	[3](S) 57		-36.8°				0.74-1.90(13H, m), 1.44(9H, s), 3.18(2H, m), 3.71(1H, m), 4.20(1H, m), 4.75(1H, d, J=10Hz), 7.22-7.59(4H, m)
15	m-cyanophenyl	70	2.8	68	114~ 117	-14.7°	C ₂₁ H ₁₃ N ₁ O ₄	C: 70.88 H: 8.24 N: 6.36	C: 70.87 H: 8.27 N: 6.16	2236, 1693, 1602, 1479, 1450, 1394, 1369, 1172, 1088
16	p-methyl-sulfonyl-aminophenyl	67	1.5	51	131~ 132	-3.3°	C ₂₁ H ₁₅ N ₁ O ₂ S	C: 61.39 H: 7.93 N: 5.51 S: 6.30	C: 61.00 H: 7.85 N: 5.48 S: 6.22	1691, 1656, 1607, 1578, 1496, 1453, 1394, 1369, 1342, 1279, 1156, 1089, 967, 918

Table 1 (continued)

Compd. of Prep. No.	R ¹	[3]		Yield%	mp°C	[α] _D ²⁰ C=1.0, CHCl ₃ (°C)	Elemental analysis	Calcd.	Found	IR (ν _{max} , cm ⁻¹)
		Yield%	C-2 S/R							
17	p-trifluoro- methylphenyl	75	4.9	80	128~ 130	-1.6 (24)	C ₉ H ₆ F ₃ NO ₄	C: 64.58 H: 7.50 N: 2.90 F: 11.79	C: 64.83 H: 7.54 N: 2.89 F: 12.02	1690, 1582, 1510, 1450, 1325, 1172, 1137, 1066
18	m-morpholino- carbonylphenyl	70	2.7	80	154~ 157	-3.6 (23)	C ₉ H ₈ N ₂ O ₄	C: 68.16 H: 8.39 N: 5.30	C: 68.04 H: 8.44 N: 5.36	1690, 1632, 1484, 1451, 1438, 1394, 1369, 1303, 1277, 1172, 1141, 1116, 1087, 1025
19	m-(N- morpholino)- ethoxyphenyl	71	3.0	39	117~ 119	-13.5 (23.5)	C ₁₃ H ₁₄ N ₂ O ₄ · 1/4 H ₂ O	C: 67.33 H: 8.90 N: 5.10	C: 67.55 H: 8.72 N: 5.06	1686, 1650, 1582
20	m-(N-2-formyl)- methylamino- phenyl	85	2.8	53	117~ 118	-16.5 (24.0)	C ₁₁ H ₁₂ N ₂ O ₃	C: 68.62 H: 8.53 N: 5.93	C: 68.68 H: 8.43 N: 5.93	1680, 1602, 1585, 1486, 1476, 1447, 1393, 1378, 1367

Preparation 21

Boc-His(Ts)-1(S)-cyclohexylmethyl-2(S)-hydroxy-4-oxo-4-(4-pyridyl)butylamide [10a]



To a 3-Boc-4-(S)-cyclohexylmethyl-2,2-dimethyl-5(S)-[2-oxo-2-(4-pyridyl)ethyl]oxazolidine[4a](4.66g, 11.18mmol) is dissolved in ethanol (20ml) is added sodium borohydride (508mg, 13.42mmol) with stirring and

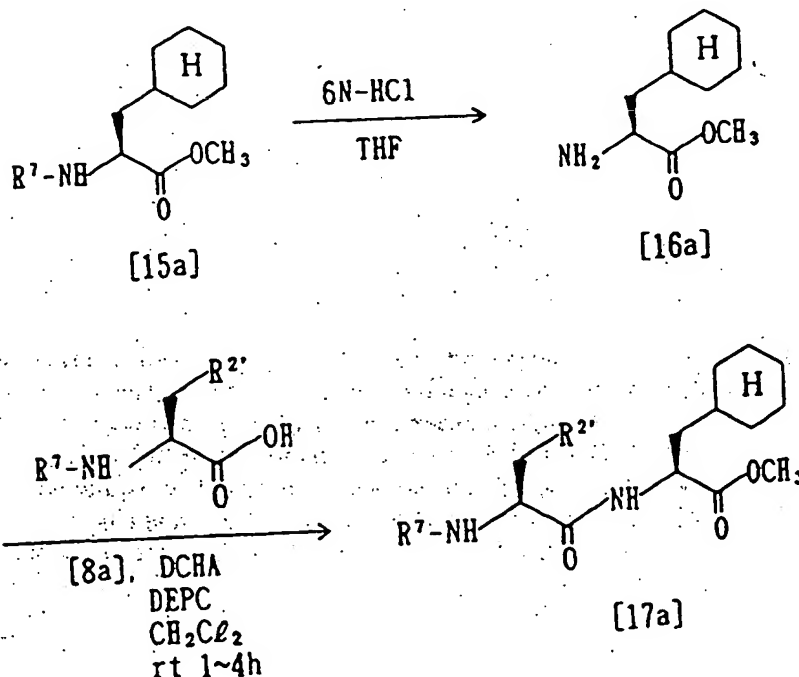
ice-cooling and the mixture is allowed to react at room temperature for one hour. The solvent is removed in vacuo. To the residue are added ice water and saturated aqueous ammonium chloride, and the mixture is extracted with dichloromethane three times. The organic layer is washed with saturated aqueous sodium chloride, dried over MgSO_4 , and evaporated to dryness in vacuo to obtain 3-Boc-4(S)-cyclohexylmethyl-2,2-dimethyl-5(S)-[2-hydroxy-2-(4-pyridyl)ethyl]oxazolidine [6a] (4.88g, quantitative amount) in colorless powder. The product is then, without further purification, dissolved in THF (2ml), and 6N HCl (16ml) is added thereto, and the mixture is stirred at room temperature for one hour. The reaction mixture is neutralized with 6N NaOH, alkalified with sodium bicarbonate, and then extracted five times with dichloromethane containing 10% methanol. The extract is dried over MgSO_4 and evaporated to dryness in vacuo to obtain 1(S)-cyclohexylmethyl-2(S), 4-dihydroxy-4-(4-pyridyl)butylamine [7a] (3.3g, quantitative amount, diastereomer ratio 1:1) in colorless powder. The product (3.30g) is then, without further purification, dissolved in dichloromethane (100ml). To the solution are added Boc-His(Ts).DCHA [8a] (8.3g, 14.05mmol, 1.3eq) and diethyl cyanophosphonate (2.29g, 14.05mmol, 1.3eq), and the mixture is stirred for 6 hours at room temperature. The reaction mixture is evaporated to dryness in vacuo, and the residue is purified with silica gel chromatography ($\text{CH}_2\text{Cl}_2:\text{MeOH} = 95:5$) to obtain Boc-His(Ts)-1(S)-cyclohexylmethyl-2(S), 4-dihydroxy-4-(4-pyridyl)butylamide [9a] (6.00g, 80%) as a mixture of two diastereomers. The product [9a] may be used in the following reaction without separation of the two isomers.

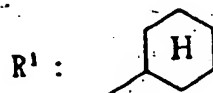
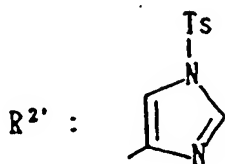
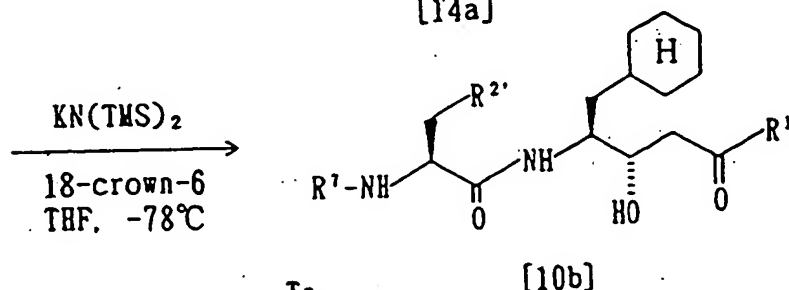
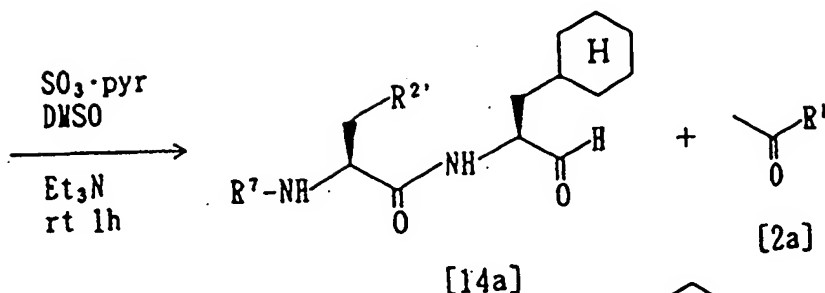
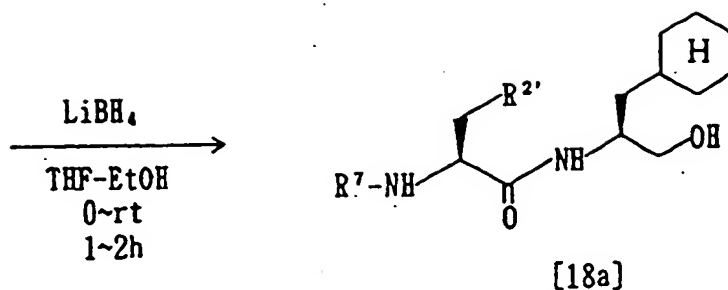
To the solution of product [9a] (1.0g, 1.45mmol) in dichloromethane (3ml) is added MnO_2 (5g) at room temperature, and the mixture is stirred for six hours. The resultant black suspension is filtered on a Celite layer overlaid with active carbon, and insoluble material on the layer is thoroughly washed with $\text{CH}_2\text{Cl}_2:\text{MeOH} (10:1)$. The filtrate is evaporated to dryness in vacuo and purified with silica gel chromatography ($\text{CH}_2\text{Cl}_2:\text{MeOH} = 95:5$) to obtain the title compound [10a] (683mg, 69%) in colorless powder.

NMR $\delta(\text{CDCl}_3)$: 1.34(9H,s), 0.70-2.20(13H,m), 2.45(3H,s), 2.99(2H,m), 3.03(1H,dd,J=17.8,2.3Hz), 3.34(1H,dd,J=17.8,9.6Hz), 4.04(1H,ddd,J=8.7,8.7,8.7Hz), 4.23(1H,m), 4.30(1H,ddd,J=5.8,5.8,5.8Hz), 6.16(1H,m), 6.47(1H,d,J=10Hz), 7.11(1H,s), 7.36(2H,d,J=8Hz), 7.80(2H,m), 7.81(2H,d,J=8.6Hz), 7.92(1H,s), 8.82(2H,d,J=5Hz)

IR $\nu(\text{CHCl}_3)$ max cm^{-1} : 3680, 3420, 3300(br), 1700, 1670, 1625, 1598, 1555, 1492, 1450, 1410, 1385, 1370, 1180, 1080, 1010

Preparation 22





$R^7 : \text{Boc}$

A solution of N-Boc-3-cyclohexyl-alanine methyl ester [15a] (4.00g, 13.93mmol) in THF (10ml) is stirred in the presence of 6N HCl (40ml) at room temperature for four hours. The reaction mixture is made alkaline with powdery sodium bicarbonate and extracted with dichloromethane containing 5% methanol (100 ml x 4).

The extract is dried over MgSO_4 and evaporated to dryness *in vacuo* to quantitatively obtain 3-cyclohexylalanine methyl ester [16a] as an oil. The product is then, without further purification, dissolved in dichloromethane (50ml). To the solution are added Boc-His(Ts).DCHA [8a] (10.7g, 18.11mmol, 1.3eq) and diethyl cyanophosphonate (2.95g, 18.1mmol, 1.3eq), and the mixture is stirred for 1.5 hours at room temperature. The reaction mixture is subjected to silica gel chromatography (SiO_2 :300g, CH_2Cl_2 :MeOH = 99:1) to give a purified Boc-His(Ts)-3-cyclohexylalanine methyl ester [17a] (7.43g, 93%) as an oil. To a solution of the dipeptide ester [17a] (3.0g, 5.2mmol) in THF (6ml) and ethanol (6ml) is added a 2N solution of lithium borohydride in THF (3ml, 6mmol) with stirring and ice-cooling. After 20 minutes stirring, the mixture is allowed to react at room temperature for an additional one hour. The solvent is removed *in vacuo* and to the residue is added ice water and saturated aqueous ammonium chloride followed by extraction with dichloromethane (20ml x 3). The organic layer is washed with saturated aqueous sodium chloride, dried over MgSO_4 , evaporated to dryness *in vacuo* and the residue is purified by silica gel chromatography (SiO_2 : 200g, CH_2Cl_2 :MeOH = 98:2) to obtain Boc-His(Ts)-3-cyclohexyl-alaninol [18a] (2.06g, 72%) as an oil.

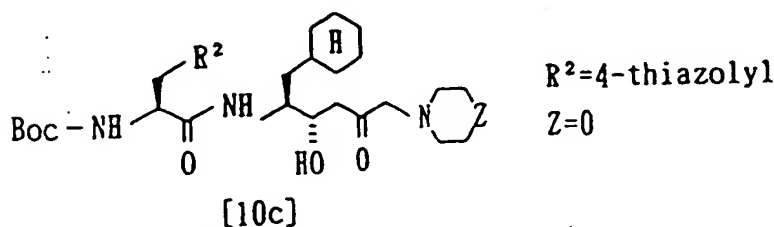
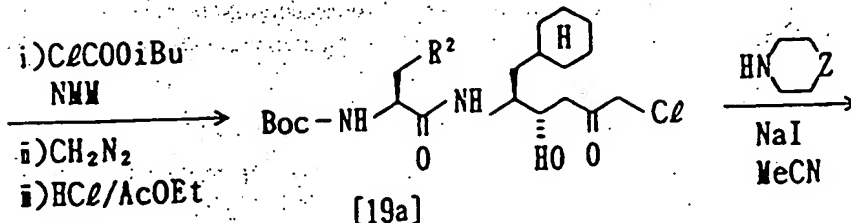
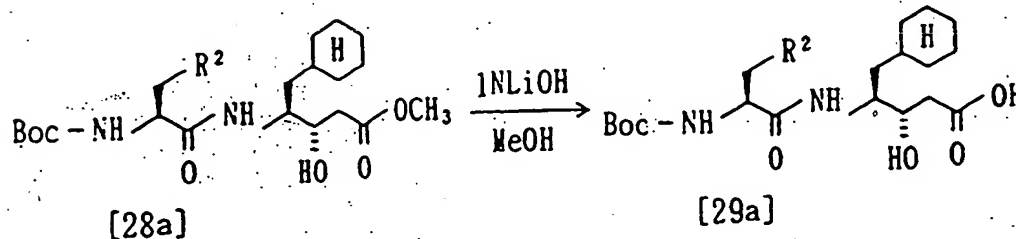
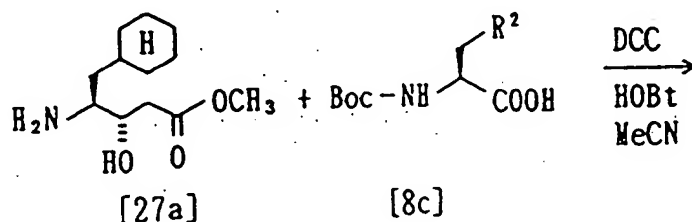
To a mixture of the dipeptide alcohol [18a] (2.0g, 3.65mmol), triethylamine (1.30g, 12.85mmol, 3.5eq) and

DMSO (6ml) is added at room temperature SO_3 -pyridine (2.03g, 12.75mmol, 3.5eq) in DMSO (6ml) and the mixture is stirred for 35 minutes. The reaction mixture is poured on ice, and the resultant aqueous mixture is extracted with ethyl acetate (20ml x 3). The organic layer is subsequently washed with 10% aqueous citric acid, saturated aqueous sodium chloride (x 2), 7% aqueous sodium bicarbonate, and saturated aqueous sodium chloride (x 2), dried over MgSO_4 , and concentrated to dryness *in vacuo*. The resultant residue is purified with silica gel chromatography (SiO_2 :100g, CH_2Cl_2 :MeOH = 95:5) to obtain Boc-His(Ts)-3-cyclohexylalaninal [14a] (1.67g, 84%) in amorphous powder.

To a 0.5N potassium bis-trimethylsilylamide solution in toluene (9.2ml, 4.60mmol, 2.5eq) is added dropwise at -78°C cyclohexyl methyl ketone (0.58g, 4.60mmol, 2.5eq) in THF (9ml) with stirring under a nitrogen atmosphere over 10 minutes. After 20 minutes stirring at the same temperature, 18-crown-6 (1.216g, 4.60mmol, 2.5eq) in THF (10ml) is dropwise added to the mixture over two minutes. Further, the dipeptidealdehyde [14a] (1.0g, 1.83mmol) in THF (10ml) is dropwise added over 15 minutes at -78°C , and the mixture is stirred for one hour at the same temperature. The reaction is quenched by adding a solution of acetic acid (0.60g, 10mmol, 5.5eq) in THF (10ml) and after the addition of saturated aqueous ammonium chloride (30ml) the mixture is extracted with ethyl acetate (50ml x 3). The organic layer is washed with saturated aqueous sodium chloride, dried over MgSO_4 , concentrated to dryness *in vacuo*, and purified with silica gel chromatography (Lobar column, CH_2Cl_2 :MeOH = 95:5) to obtain Boc-His(Ts)-1(S)-cyclohexylmethyl-2(S)-hydroxy-4-oxo-4-cyclohexyl-butylamide [10b] (0.18g, 15%) in amorphous powder.

NMR δ : 1.30-1.90(23H,m), 1.40(9H,s), 2.32(1H,m), 2.44(3H,s), 2.59(2H,m), 2.93(1H,dd,J=5.8,9.6Hz), 3.04(1H,dd,J=5.8,9.6Hz), 3.89(1H,ddd,J=8.4,8.4,8.4Hz), 3.98(1H,m), 4.30(1H,ddd,J=6.0,6.0,6.0Hz), 6.12(1H,d,J=6.0Hz), 6.47(1H,d,J=9.8Hz), 7.10(1H,d,J=0.8Hz), 7.36(2H,d,J=8.0Hz), 7.81(2H,d,J=8.4Hz), 7.93(1H,d,J=1.2Hz)

Preparation 23



To a solution of cyclostatine methyl ester [27a] (700mg, 3.05mmol), Boc-(4-thiazolyl)-L-alanine [8c] (869mg, 3.19mmol, 1.05eq), and HOBt (431mg, 3.19mmol, 1.05eq) in CH₃CN (10ml) is added DCC (660mg, 3.20mmol, 1.05eq) with stirring and ice-cooling under nitrogen atmosphere and the mixture is stirred for 1.5 hours at the same temperature and then allowed to react at room temperature for 14 hours. Ethyl acetate is added to the mixture, and precipitated crystals were filtered off. The filtrate is concentrated to dryness in vacuo and the residue is subjected to silica gel chromatography (SiO₂:100g, NH₄OH:MeOH:CH₂Cl₂ = 1:10:990) to give the aimed product, Boc-(4-thiazolyl)alanyl-cyclostatine methyl ester [28a] (830mg, 59%) as an oil.

To the solution of the above product [28a] (830mg, 1.72mmol) in MeOH (2ml) is added 1N LiOH (1.9ml, 1.9mmol, 1.1eq) with stirring and ice-cooling. The mixture is stirred for 10 minutes and allowed to react at room temperature for two hours. After neutral substances are removed by washing with dichloromethane, the mixture is acidified with citric acid and is extracted with ethyl acetate. The organic layer is dried over MgSO₄, and concentrated to dryness in vacuo to obtain the aimed carboxylic acid [29a] (700mg, 87%).

To a mixture of the above carboxylic acid [29a] (700mg, 1.67mmol) and N-methylmorpholine (0.17ml, 1.67mmol) in THF (10ml) is added isobutyl chlorocarbonate (0.2ml, 1.67mmol) with stirring at a temperature of -15°C - -10°C under nitrogen atmosphere, and the resultant mixture is stirred for 50 minutes at the same temperature. After precipitated crystals are removed by filtration, to the filtrate is added a solution of diazomethane (2.2eq) in ethyl ether previously prepared at -10°C and allowed to react at room temperature for 3 hours. The reaction mixture is concentrated in vacuo to remove diazomethane and ethyl acetate (10ml) is added to the residue. After addition of 2N HCl (3ml) at -40°C - -30°C, the mixture is allowed to react for one hour. The reaction mixture is alkalified by addition of saturated aqueous sodium bicarbonate and the ethyl acetate layer is separated. The layer is dried over MgSO₄ and concentrated to dryness in vacuo to obtain 800mg of crude chloromethyl ketone [19a]. Since the product tends to get colored and decomposed, it is immediately used in the next step without purification.

To a solution of the above product [19a] (400mg) in MeCN (5ml) are added morpholine (150mg) and a catalytic amount of NaI, and the mixture is stirred at room temperature for two hours. The reaction mixture is purified by chromatography to give the aimed compound, Boc-(4-thiazolyl)alanyl-1(S)-cyclohexylmethyl-2(S)-hydroxy-4-oxo-4-(N-morpholino)methyl-butylamide [10c] (Z=O) (120mg, 29% starting from [29a]).

NMRδ: 0.6-2.00(13H,m), 1.43(9H,s), 2.55(4H,m), 3.22(2H,dd,J=4.6,14.8Hz), 3.26(2H,s), 3.43(1H,dd, J=5.4,14.8Hz), 3.76(4H,m), 3.89(1H,m), 3.94(1H,m), 4.44(1H,ddd,J=6.2HzX3), 6.38(1H,d,J=9.8Hz), 6.48 (1H, d, J=7.5Hz), 7.13(1H,d,J=1.8Hz), 8.79(1H,d,J=2Hz)

Preparation 24

In the same manner as in Preparation 23, Boc-(4-thiazolyl)alanyl-1(S)-cyclohexylmethyl-2(S)-hydroxy-4-oxo-4-(N-piperidino)methyl-butylamide [10d] (Z=CH₂) is obtained with an overall yield of 29%.

NMRδ: 0.6-1.83(19H,m), 1.44(9H,s), 2.46(4H,m), 3.15(2H,s), 3.20(1H,dd,J=5.6,14.8Hz), 3.44(1H, dd, J=5,14.8Hz), 3.89(2H,m), 4.47(1H,m), 6.41(1H,bs), 6.43(1H,d,J=9.8Hz), 7.12(1H,d,J=1.8Hz), 8.78 (1H, d, J=1.8Hz)

Preparation 25-50

Starting from the compounds [4] which have been prepared in Preparations 2-20, the ketone compounds [10] are obtained in the same manner as in Preparation 21. The thus obtained products are listed in Table 2.

Preparation 51-57

The aldol reaction between dipeptides [14] and methyl ketones [2] gives ketone compounds [10] in the same manner as in Preparation 22. The thus obtained products are listed in Table 3.

Table 2 (continued)






Compd. of Prep. No.	R'	R	[9]		[10]
			Yield%	Yield% (from 7)	
28	p-ethylphenyl		78	57	3420, 3300, 3240, 1705, 1670, 1625, 1608, 1495, 1450, 1370, 1122, 1033, 1010
29	2,4-difluorophenyl		72	50	3680, 3420, 3300(br), 1705, 1672, 1611, 1599, 1496, 1450, 1430, 1384, 1370, 1172, 1095, 1080, 970, 855
30	1-naphthyl		71	50	3692, 3420, 1709, 1673, 1599, 1575, 1495, 1450, 1386, 1370, 1175, 1094, 1080, 1033, 979, 908
31	3-thienyl		48	63	0, 7-1, 85(13II, s), 1, 34(9II, s), 2, 44(3II, s), 2, 99(2II, m), 2, 86-3, 27(2II, m), 4, 00(1II, ddd, J=9, 9, 9IIz), 4, 19(1II, d, J=10IIz), 4, 32(1II, ddd, J=6, 6, 6IIz), 6, 15(1II, d, J=5, 0IIz), 6, 54 (1II, d, J=9IIz), 7, 11(1II, d, J=1, 2IIz), 7, 36(2II, d, J=8, 2IIz), 7, 3(1II, dd, J=2, 8, 5, 2IIz), 7, 53(1II, dd, J=1, 2, 5, 2IIz), 7, 80(2II, d, J=8, 4IIz), 7, 91(1II, d, J=1, 2IIz), 8, 18(1II, s)
32	2-thiazolyl		79	5	3420, 3300, 3140, 1703, 1670, 1625, 1603, 1550(br), 1496, 1450, 1370, 1165, 1123, 1032, 1010

Table 2 (continued)









Compd. of Prep. No.	R ¹	R ²	[9]		[10]	
			Yield% (from [7])	Yield%	IR ν _{max} cm ⁻¹ , NMR(δ), [α] _D ²⁰ , m.p.	
33	m-fluorophenyl	Ts 	76	45	3420, 3280, 3140, 1675, 1625, 1590, 1495, 1160, 1122, 1030, 1010	
34	p-fluorophenyl	Ts 	78	61	3420, 3320, 3140, 1670(shoulder), 1625, 1600, 1495, 1155, 1030, 1010	
35	2,6-di-fluorophenyl	Ts 	88	30	3424, 1707, 1676, 1625, 1598, 1495, 1468, 1386, 1370, 1174, 1094, 1080, 1028 [α] _D ²⁰ = -6.0° (C=1.0, CHCl ₃ , 24°C)	
36	o-acethoxyphenyl	Ts 	31	55	0.75-1.83(13H, m), 1.36(9H, s), 2.44(3H, s), 3.00(2H, m), 3.13(2H, m), 3.90(3H, s), 3.96(1H, ddd, J=10.10, 10Hz), 4.15(1H, m), 4.34(1H, ddd, J=7.7, 7Hz), 6.04(1H, d, J=7Hz), 6.58(1H, d, J=10Hz), 6.99(2H, m), 7.10(1H, d, J=1.2Hz), 7.35(2H, d, J=8.6Hz), 7.47(1H, m), 7.70(1H, dd, J=2.7, 8Hz), 7.80(2H, d, J=8.4Hz), 7.91(1H, d, J=1.4Hz)	
37	o-chlorophenyl	Ts 	85	27	0.74-1.82(13H, m), 1.39(9H, s), 2.44(3H, s), 2.98(2H, m), 3.08(2H, m), 3.99(1H, m), 4.18 (1H, m), 4.30(1H, ddd, J=7Hz), 6.05(1H, d, J=7Hz), 6.52(1H, d, J=10Hz), 7.10(1H, d, J=1.3Hz), 7.27-7.45(6H, m), 7.60(1H, m), 7.80(2H, d, J=8.4Hz), 7.90(1H, d, J=1.4Hz)	
38	m-cyanophenyl	Ts 	67	51	3420, 2236, 1709, 1678, 1599, 1495, 1451, 1432, 1387, 1371, 1174, 1093, 1081, 909	
39	o-ethyl-sulfonyl-aminophenyl	Ts 	76	58	3424, 1709, 1672, 1599, 1578, 1496, 1452, 1385, 1371, 1341, 1174, 1155, 1094, 1081, 1034, 968, 917	
40	p-trifluoroethyl-phenyl	Ts 	74	37	3420, 1705, 1675, 1625, 1575, 1325, 1170, 1135, 1080, 1065 m.p. = 133-135°C	

Table 2 (continued)








Compd. of Prep. No.	R ¹	R ²	[9]		[10]	
			Yield% (from [7])	Yield%	I R, ν_{max} cm ⁻¹ またはNMR (δ)	
41	m-morpholino- carbonylphenyl	Ts 	85	68		3420, 1709, 1674, 1632, 1600, 1495, 1386, 1370, 1279, 1174, 1116, 1093, 1080, 1026
42	phenyl		90	57		0. 70~1. 90(13H, m), 1. 34(9H, s), 2. 90~3. 60(4H, m), 3. 99(1H, m), 4. 16(1H, m), 4. 49(1H, ddd, J=6. 2, 6. 2, 6. 2 Hz), 6. 46(1H, d, J=9. 2 Hz), 7. 12(1H, d, J=1. 8 Hz), 7. 40~7. 63(3H, m), 7. 96(2H, d, J=8. 4 Hz), 8. 76(1H, d, J=2 Hz)
43	4-pyridyl		90	64		0. 70~2. 05(13H, m), 1. 34(9H, s), 2. 95~3. 50(4H, m), 4. 01(1H, m), 4. 19(1H, m), 4. 46(1H, ddd, J=5. 8 Hz), 6. 4(1H, d, J=8 Hz), 6. 55(1H, d, J=5 Hz), 7. 13(1H, d, J=1. 8 Hz), 7. 78(2H, d, J=6. 2 Hz), 8. 77(1H, d, J=1. 8 Hz), 8. 80(1H, d, J=9. 4 Hz)
44	3-thienyl		86	80		0. 65~2. 05(13H, m), 1. 36(9H, s), 2. 93(1H, d, J=17. 1 Hz), 3. 15(1H, dd, J=17. 6, 9. 4 Hz), 3. 22(1H, dd, J=14. 8, 5. 4 Hz), 3. 44(1H, dd, J=14. 8, 5. 3 Hz), 3. 97(1H, m), 4. 15(1H, m), 4. 48(1H, ddd, J=6. 4, 6. 4, 6. 4 Hz), 6. 44(1H, d, J=9. 9 Hz), 6. 50(1H, d, J=7. 5 Hz), 7. 12 (1H, d, J=1. 9 Hz), 7. 30(1H, dd, J=5. 1, 2. 9 Hz), 7. 53(1H, dd, J=5. 1, 1. 3 Hz), 8. 20(1H, d, J=1. 9 Hz), 8. 77(1H, d, J=2. 1 Hz)
45	m-2-(N- morpholino)- ethoxyphenyl		96	51		0. 70~1. 90(13H, m), 1. 35(9H, s), 2. 60(4H, m), 2. 83(2H, t, J=5. 4 Hz), 3. 17(2H, m), 3. 22(1H, dd, J=4. 6, 14. 4 Hz), 3. 44(1H, dd, J=5. 2, 14. 6 Hz), 3. 75(4H, m), 3. 98(1H, ddd, J=6. 2, 6. 2, 6. 2 Hz), 4. 49(3H, m), 4. 49(1H, ddd, J=6. 2, 6. 2, 6. 2 Hz), 6. 45(2H, d, J=9. 8 Hz), 7. 11(1H, d, J=2. 0), 7. 15(1H, dd, J=1. 2, 2. 8 Hz), 7. 37(1H, t, J=7. 8 Hz), 7. 48(1H, m), 7. 56(1H, d, J=7. 8 Hz), 8. 77(1H, d, J=2. 0 Hz)
46	m-(N-formyl)- methylamino		97	77		0. 70~1. 86(13H, m), 1. 33(9H, s), 3. 00(1H, dd, J=0. 5, 14. 8 Hz), 3. 23(1H, dd, J=5. 2, 14. 8 Hz), 3. 36(3H, s), 3. 40(3H, m), 4. 01(1H, m), 4. 20(1H, d, J=9. 8 Hz), 4. 47(1H, ddd, J=5 Hz), 6. 44(1H, d, J=9. 8 Hz), 6. 58(1H, d, J=6. 4 Hz), 7. 14(1H, d, J=1. 8 Hz), 7. 39(1H, ddd, d, J=1. 2, 2. 4, 8 Hz), 7. 52(1H, t, J=7. 8 Hz), 7. 87(2H, m), 8. 57(1H, s), 8. 78(1H, d, J=2 Hz)
47	4-pyridyl		87	61		0. 77~1. 84(13H, m), 2. 70(3H, s), 3. 17(4H, m), 4. 03(1H, m), 4. 20(1H, m), 4. 42(1H, ddd, J=5. 8 Hz x 3), 6. 42(1H, d, J=5 Hz), 6. 49(1H, d, J=10 Hz), 6. 89(1H, s), 7. 80(2H, m), 8. 81(2H, m)

Table 2 (continued)


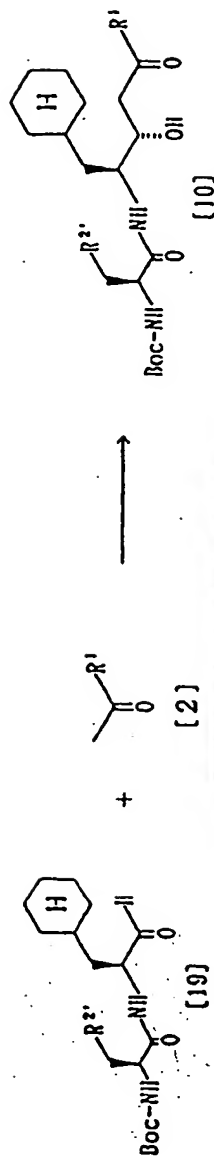
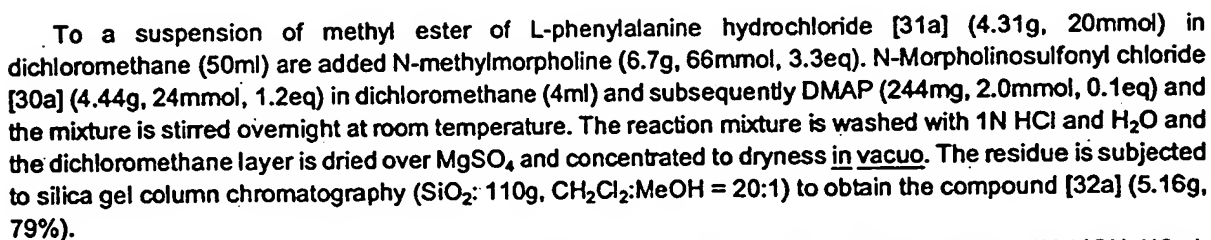
Compd. of Prep. No.	R ¹	R ²	[9]		[10]	
			Yield% (from [7])	Yield%	IR ν_{\max} cm ⁻¹ or NMR (δ)	
48	phenyl		62	89	0.77~1.82(13H, m), 1.38(9H, s), 3.10(4H, m), 4.03(1H, m), 4.16(1H, m), 4.41(1H, t, J=5.2Hz), 6.71(1H, s), 6.89(1H, d, J=8.2Hz), 7.47(2H, t, J=7.8Hz), 7.59(1H, m), 7.94(2H, d, J=7.2Hz), 8.47(1H, s)	
49	4-pyridyl	-CONH ₂	86	65	0.76~1.82(13H, m), 1.41(9H, s), 2.67(2H, m), 3.06(1H, dd, J=4.2, 18Hz), 3.27(1H, dd, J=8.4, 18Hz), 4.04(1H, m), 4.24(1H, m), 4.37(1H, t, J=6.4Hz), 7.07(1H, d, J=9.4Hz), 7.85(2H, m), 8.76(2H, m)	
50	4-pyridyl	-SMe	66	36	0.82~1.88(13H, m), 1.38(9H, s), 2.15(3H, s), 2.87(1H, dd, J=6.4, 13.6Hz), 2.95(1H, t, J=7.6Hz), 3.10(1H, dd, J=2.2, 18.8Hz), 3.42(1H, dd, J=9.4, 18.6Hz), 4.12(1H, m), 4.23(1H, ddd, J=6Hzx3), 4.28(1H, m), 5.37(1H, d, J=6Hz), 6.55(1H, d, J=10Hz), 7.77(2H, m), 8.82(2H, m)	

Table 3



Compd. of Prep. No.	R ¹	R ²	Yield%	NMR δ (CDCl ₃)
51	p-acethoxyphenyl	TS	10	0.75-1.94 (13H, m), 1.33 (9H, s), 2.44 (3H, s), 3.00 (2H, m), 3.88 (3H, s), 4.01 (1H, ddd, J=8.2 Hz), 4.19 (1H, m), 4.34 (1H, ddd, J=6.4 Hz), 6.12 (1H, d, J=5.8 Hz), 6.58 (1H, d, J=9.8 Hz), 6.93 (2H, d, J=8.6 Hz), 7.11 (1H, s), 7.36 (2H, d, J=8.2 Hz), 7.81 (2H, d, J=8.6 Hz), 7.93 (1H, s), 7.95 (2H, d, J=9.1 Hz), 7.77-1.83 (13H, m), 1.34 (9H, s), 2.44 (3H, s), 3.00 (4H, m), 4.00 (1H, ddd, J=8.4, 8.4, 8.4 Hz), 4.18 (1H, d, J=6.2 Hz), 4.32 (1H, ddd, J=6.6 Hz), 6.05 (2H, s), 6.13 (1H, m), 6.54 (1H, d, J=9.8 Hz), 6.85 (1H, d, J=8.2 Hz), 7.11 (1H, d, J=0.4 Hz), 7.36 (2H, d, J=8.4 Hz), 7.43 (1H, d, J=1.4 Hz), 7.58 (1H, td, J=8.2, 0.8 Hz), 7.80 (2H, d, J=8.4 Hz), 7.92 (1H, d, J=1.2 Hz)
52	3,4'-acetylendioxyphenyl	TS	19	Identical with those of compound in Ex. No. 27
53	3-thienyl	TS	23	Identical with those of compound in Ex. No. 27
54	morpholinocarbonyloxyphenyl	TS	27	0.72-2.00 (13H, m), 1.34 (9H, s), 2.44 (3H, s), 3.00 (2H, m), 3.52-3.8 (8H, m), 4.00 (1H, ddd, J=8.1 Hz), 4.18 (1H, m), 4.33 (1H, ddd, J=6.6 Hz), 6.10 (1H, m), 6.58 (1H, d, J=7.1 Hz), 7.12 (1H, d, J=3.4 Hz), 7.36 (3H, m), 7.47 (1H, t, J=8.1 Hz), 7.68 (1H, m), 7.82 (3H, m), 7.94 (1H, d, J=1.4 Hz)
55	phenyl	TS	28	Identical with those of compound in Ex. No. 21
56	N-acetyl-3-pyrrolyl	TS	39	0.70-1.85 (13H, m), 1.38 (9H, s), 2.81 (2H, d, J=5.6 Hz), 3.20 (1H, dd, J=4.8, 14.2 Hz), 3.45 (1H, dd, J=5.2, 14.8 Hz), 3.69 (3H, s), 3.92 (1H, m), 4.07 (1H, t, J=5.8 Hz), 4.49 (1H, ddd, J=6.8 Hz), 4.93 (1H, d, J=9.6 Hz), 6.57 (1H, s), 6.58 (1H, s), 7.12 (1H, d, J=2.1 Hz), 7.32 (1H, s), 8.77 (1H, d, J=2.1 Hz), 7.6-1.92 (13H, m), 2.33 (1H, m), 2.45-2.75 (2H, m), 3.20 (1H, dd, J=14.4, 5.2 Hz), 3.44 (1H, dd, J=14.8, 3.8 Hz), 3.85 (1H, m), 3.93 (1H, m), 4.45 (1H, ddd, J=6.2, 6.2, 2.1 Hz), 6.40 (1H, d, J=9.8 Hz), 6.49 (1H, d, J=6.8 Hz), 7.12 (1H, d, J=1.6 Hz), 8.78 (1H, d, J=1.8 Hz)
57	cyclohexyl	TS	31	Identical with those of compound in Ex. No. 21



(i) To a solution of the compound [32a] (2.666g, 8.1mmol) in MeOH (12ml) is added 1N LiOH (12ml, 12mmol, 1.5eq) and the mixture is stirred at 80°C for 30 minutes. After removal of MeOH in vacuo, the reaction mixture is washed with ethyl acetate. The mixture is then treated with active carbon, adjusted to pH 2 - 3 with 1N HCl, and extracted with ethyl acetate. The extract is washed with saturated aqueous sodium chloride, dried over MgSO₄, and concentrated to dryness in vacuo. The residue is recrystallized from ethyl acetate/n-hexane to colorless needles of N-(N-morpholinosulfonyl)-phenylalanine [12b] (2.267g, 89%). m.p. 164 - 6°C (decomposition)

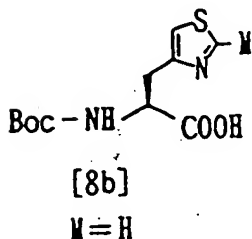
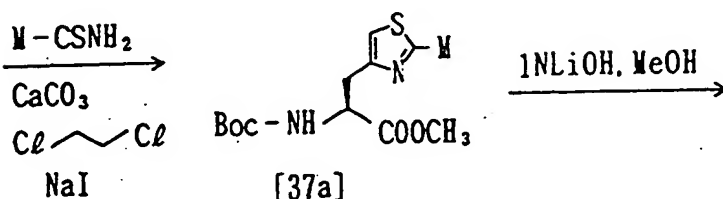
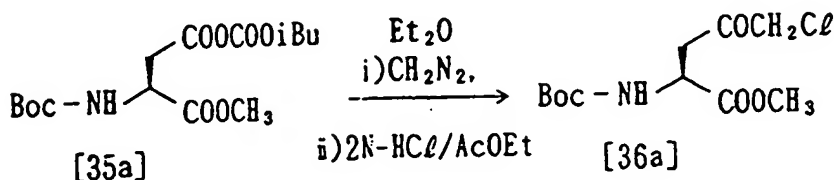
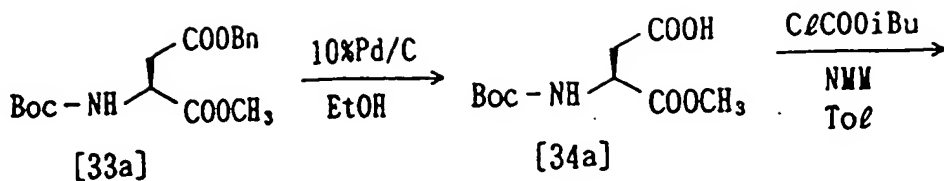
(ii) To the compound [32b] (E=Et) (920mg, 2.7mmol) are added 6N HCl (9.2ml) and acetic acid (2ml) and the mixture is heated with stirring on an oil bath of 100°C for one hour. After cooling, the reaction mixture is concentrated to dryness in vacuo. The residue is made alkaline by dissolving into saturated aqueous sodium bicarbonate. The aqueous solution is washed with dichloromethane (10ml x 3), treated with active carbon, and neutralized with 6N HCl. The solution is then made acidic up to pH 3 by addition of 10% aqueous citric acid and extracted with ethyl acetate (50ml x 3). The organic layer is washed with saturated aqueous sodium chloride (x 2), dried over MgSO₄, and concentrated to dryness in vacuo to give the compound [12c] as a crystalline residue (620mg, 74%). Recrystallization from dichloromethane/isopropyl ether affords white crystals (543mg, 64%). m.p. 157 - 158°C.

$$[\alpha]_D = -17.7 \pm 0.6^\circ (C=1.0; \text{MeOH}; 25.0^\circ\text{C})$$

IR_vmax (cm⁻¹): 3320, 3200-2600(br), 1750, 1603, 1585, 1500, 1455, 1400, 1352, 1300

NMR(δ): 2.93(5H,m), 3.17(1H,dd,J=5.2,14.2Hz), 3.54(4H,m), 4.11(1H,dd,J=5.2,8.6Hz), 7.30(5H,m)

Preparation 59



a) A solution of methyl ester of N-Boc- ω -benzyl-L-aspartic acid [33a] (52.7g, 0.156mmol) in a mixture of water (10ml), acetic acid (10ml) and methanol (150ml) is subjected to a catalytic reduction in the presence of 10% Pd-C (4.0g) under an atmosphere of hydrogen gas at room temperature. The reduction is conducted with stirring and under atmospheric pressure. After a 3-hour reaction, the catalyst is filtered off and the filtrate is evaporated to dryness *in vacuo*. The residue is dissolved in saturated aqueous sodium bicarbonate and the aqueous layer is washed with dichloromethane (50ml x 3), made acidic with citric acid (about pH3), and extracted with ethyl acetate (200ml x 4) while salting out with the addition of sodium chloride. The ethyl acetate layer is dried over MgSO₄ and concentrated to dryness *in vacuo*. Trituration of the residue with the addition of n-hexane affords the carboxylic acid [34a] (37.5g, 98%) as a white solid.

To a solution of the above product [34a] (18.8g, 76mmol) and N-methylmorpholine (7.8g, 77.1mmol, 1.0eq) in ethyl ether (200ml) is added isobutyl chlorocarbonate (9.92ml, 76.5mmol, 1.0eq) over 10 minutes at a temperature between -15°C and -10°C under nitrogen atmosphere, and the mixture is stirred at the same temperature for 30 minutes. Preprecipitated methylmorpholine hydrochloride is filtered off, and the filtrate is added to a solution of diazomethane in ethyl ether which has previously been prepared from nitrosomethylurea (37g, 359mmol) with stirring at -10°C over 5 minutes. After 2.5-hour stirring at room temperature, the mixture is concentrated *in vacuo* to remove excessive diazomethane. To the mixture is added ethyl acetate (150ml) and then dropwise added 2N HCl/ethyl acetate (45ml) at a temperature between -40°C and -30°C. After 30-minutes stirring, the mixture is neutralized with saturated aqueous sodium bicarbonate. The ethyl acetate layer is separated, dried over MgSO₄, evaporated to dryness *in vacuo*, and subjected to silica gel chromatography (SiO₂: 150g, AcOEt:CH₂Cl₂ = 6:1) to obtain the chloromethyl ketone [36a] (20.3g, 95%) as an oil.

To a solution of the above compound [36a] (40.3g, 144.1mmol) in MeCN (160ml) are added CaCO₃ (28g, 280mmol, 1.9eq) and thioformamide (HCSNH₂, 14g, 229.1mmol, 1.6eq) and the mixture is stirred at room temperature for 18 hours under nitrogen atmosphere. Insoluble materials are filtered off and the filtrate is concentrated to dryness *in vacuo*. The residue is dissolved in dichloromethane, subsequently washed with 7% aqueous

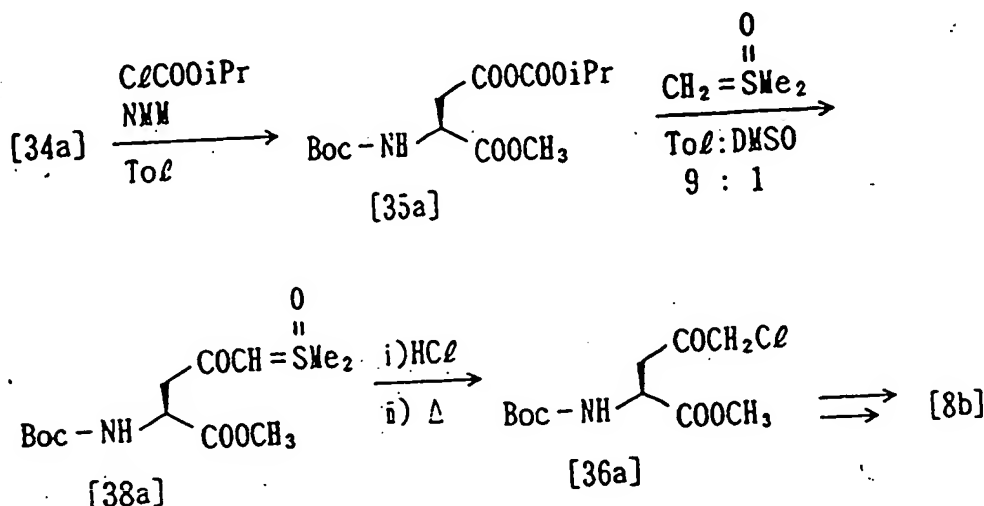
sodium bicarbonate, 1N NaOH, and water, two times each, to remove non-reacted thioformamide. The dichloromethane layer is dried over MgSO_4 , concentrated to dryness in vacuo, and subjected to silica gel chromatography (SiO_2 : 370g, $\text{MeCN}:\text{CH}_2\text{Cl}_2 = 1:7$) to obtain (4-thiazolyl)alanine derivative [37a] (29.15g, 71%) as an oil.

To the solution of above product [37a] (29.1g, 101.6mmol) in methanol(120ml) is added 1N LiOH (112ml, 112mmol, 1.1eq) with stirring and ice-cooling and the mixture is stirred for ten minutes at the same temperature and allowed to react an additional one hour at room temperature. The reaction mixture is concentrated in vacuo on a water bath below 30°C to remove methanol and the residue is washed three times with dichloromethane. The aqueous layer is treated with active carbon, and citric acid to adjust the pH to 3, and extracted with ethyl acetate (150ml x 3). To the organic layer washed two times with saturated aqueous sodium chloride are added MgSO_4 and active carbon, the mixture is filtered and the filtrate is concentrated to dryness in vacuo to obtain crystalline crude product [8b] (26.96g, 97%). Recrystallization of the product from n-hexane provides pure product [8b] (26.2g, 95%). m.p. $96 - 98^\circ\text{C}$

$[\alpha]_D^{25} = -4.2^\circ (c=2; \text{MeOH}; 24^\circ\text{C})$

NMR(δ): 1.47(9H,s), 3.41(1H,dd,J=5.6,14.6Hz), 3.56(1H,dd,J=3.4,11.0Hz), 4.59(1H,m), 3.60(1H,d,J=3.6Hz), 7.14(1H,d,J=2Hz), 8.94(1H,d,J=2Hz)

b)



i) Preparation of carbonic anhydride

To a solution of compound [34a] (500mg, 2.02mmol) and N-methylmorpholine (225mg, 2.22mmol, 1.1eq) in toluene (4ml) is added isopropyl chlorocarbonate (0.254ml, 2.22mmol, 1.1eq) with stirring at a temperature between -15°C and -10°C under nitrogen atmosphere and the mixture is stirred at the same temperature for one hour to separate out N-methylmorpholine hydrochloride.

ii) Preparation of Corey reagent (dimethylsulfoxonium methylide)

To a suspension of trimethylsulfoxonium iodide (1.024g, 4.65mmol) in toluene (9ml) and DMSO (1ml) is added potassium t-butoxide (522mg, 4.65mmol, 1.0eq) with stirring under nitrogen atmosphere, and the mixture is heated with stirring on an oil bath at $70 - 75^\circ\text{C}$ for 30 minutes. Orange crystals turn to grayish white crystals.

The carbonic anhydride solution obtained in the above step i) is charged in a dropping funnel with a cotton stopper. The solution is dropwise added to the Corey reagent prepared in the step ii) from the funnel with stirring and ice-cooling under nitrogen atmosphere over 10 minutes and the mixture is stirred at room temperature for one hour. The mixture is filtered and the filtrate is extracted with water (10ml x 3). The aqueous layer is extracted with dichloromethane (10ml x 4). Each extract is washed with water, dried over MgSO_4 , and concentrated to dryness in vacuo to obtain 600mg of crude product. Chromatography (SiO_2 : 40g, 3.5% $\text{MeOH}:\text{CH}_2\text{Cl}_2$) of the crude product gives the aimed ylide compound [38a] (554mg, 85%) as an oil.

To a solution of the ylide [38a] (3.16g, 9.83mmol) in dichloroethane (26ml) is added 2N HCl/ethyl acetate (4.92ml, 9.84mmol) with stirring at -10°C and the mixture is stirred for one hour. The mixture is warmed on an oil bath of 100°C . Although precipitates (HCl addition product) separate out after two minutes, they redissolve after 3.5 minutes. When the solution becomes turbid after 6 minutes, the solution is cooled immediately to ter-

minate the reaction and the reaction mixture is subjected to silica gel chromatography (SiO_2 : 15g, $\text{AcOEt}:\text{CH}_2\text{Cl}_2$ = 1:7) to obtain chloromethyl ketone [36a] (2.308g, 84%) as a crystal substance.

A suspension of the above product [36a] (2.308g, 8.25mmol), HCSNH_2 (1.26g, 20.62mmol, 2.5eq) and CaCO_3 (2.475g, 24.75mmol, 3eq) in dichloroethane (23ml), is stirred at room temperature for 15 hours under nitrogen atmosphere. After addition of NaI (62mg, 0.414mmol, 0.05eq), the mixture is stirred for an additional two hours. Insoluble materials are filtered off and washed with dichloromethane. The filtrate and washings are combined and subsequently washed with saturated aqueous sodium bicarbonate, 1N NaOH , and H_2O (x 2). Chromatographic treatment of the solution in the same manner as described in the foregoing process a) provides (4-thiazolyl)-L-alanine derivative [37a] (1.878g, 80%) as an oil.

To a solution of the above compound [37a] (3.16g, 11.04mmol) in methanol (6ml) is added with stirring and ice-cooling 1N LiOH (13ml, 13mmol, 1.18eq) and the mixture is stirred at room temperature for one hour. Similar procedure as disclosed in the process a) provides crude product [8b] (2.9g, 97%). Recrystallization of the product from ethyl ether/n-hexane gives pure product [8b] (2.6g, 88%) as colorless crystals. m.p. 110 - 112°C. $[\alpha]_D^{25} = -4.8$ (c=2.0; MeOH; 25°C)

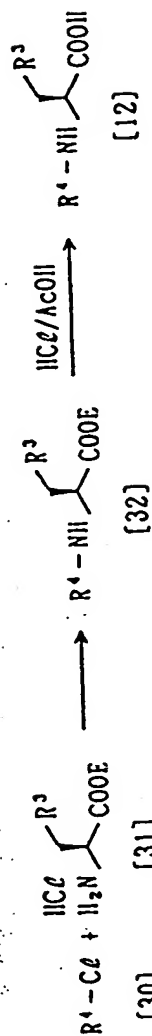
Preparation 60 and 61

N-sulfamylamino acids [12] listed in Table 4 are prepared from the compounds [30] in the same manner as disclosed in Preparation 58.

Preparation 62 and 63

2-Substituted (4-thiazolyl)-L-alanines [8] listed in Table 5 are prepared from the compounds [36] in the same manner as disclosed in Preparation 59.

Table 4




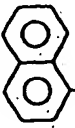
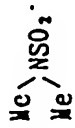

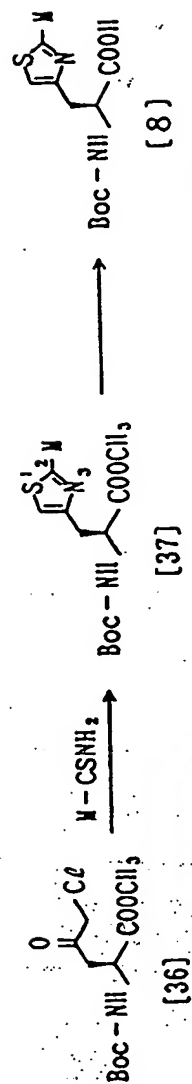
Compd. of Prp. No.	R ⁴	R ³	E	[10]			NMR (δ)
				[32] Yield%	[31] Yield%	[32] [α] _D ²⁰ (Temp. °C)	
60			Me	88	76	-56.7(25)	2.55(2H, m), 2.63(2H, m), 3.13(2H, m), 3.20(3H, m), 3.55(1H, bs), 3.80(1H, dd, J=4.6, 14Hz), 4.35(1H, dt, J=4.4, 10Hz), 5.05(1H, d, J=10.2Hz), 7.37(2H, m), 7.57(2H, m), 7.89(2H, m), 8.10(1H, d, J=8.2Hz)
61			Et	82	26		2.58(6H, s), 2.98(1H, dd, J=7.8, 13.6Hz), 3.20(1H, dd, J=5.2, 13.6Hz), 4.24(1H, dofdd, J=9.6, 7.4, 4.6Hz), 4.90(1H, d, J=10Hz), 4.90(1H, bs), 7.30(5H, m)

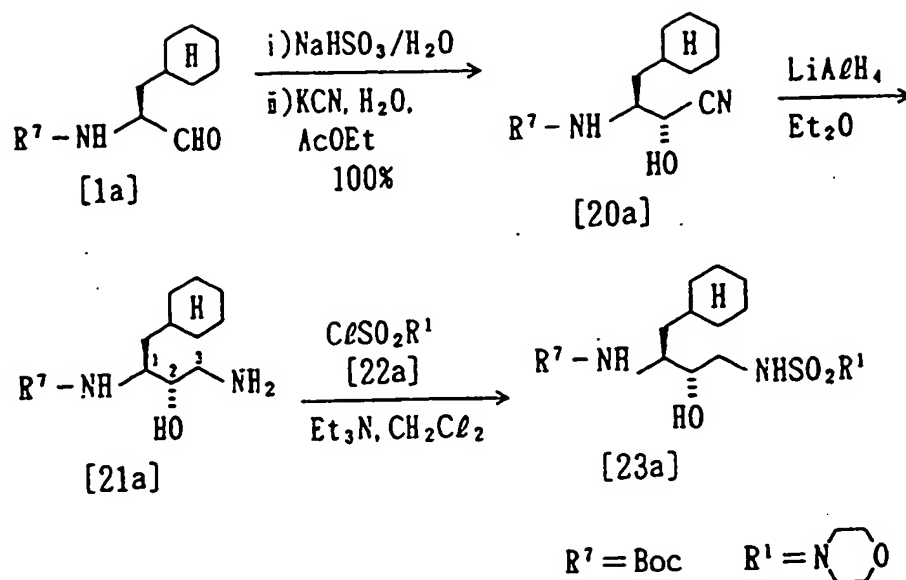
Table 5



Compd. of Prep. No.	M	[37]		mp. (°C)	Yield%	[α] _D ²⁰ C=1, MeOH (temp. °C)	IR ν _{max} cm ⁻¹	NMR (δ)
		Yield%	[α] _D ²⁰ C=1, MeOH (temp. °C)					
62	Cl ₃	42	—	135- 136	93	(C=2) -20.4 (24)	3430, 2440(br) 1700, 1495, 1435, 1392 1368, 1160 1060	
63	NH ₂	88	-345 (24) -10.1* (22)	* 156- 157	* 87	* -4.3 (22)	*3440, 3200, 2440(br) 1700, 1565, 1500 1455, 1435, 1392 1370, 1160 1062	*1.50(9H, s), 3.38(2H, m), 4.64(1H, m), 5.15(2H, d, J=6.8Hz), 6.69(1H, s), 8.48(1H, s)

* formyl compound

Preparation 64



To the aldehyde compound [1a] (10.08g, 39.5mmol) is added NaHSO₃ (10.08g) in water (70ml) and the mixture is stirred with ice-cooling for 16 hours. The resultant solution is stirred at room temperature for 4 hours after addition of KCN (6.3g) in water (16.8ml) and ethyl acetate (137ml). The ethyl acetate layer is separated from the reaction mixture, washed with saturated aqueous sodium chloride, dried, and concentrated. The residue is subjected to column chromatography using Lobar column Size C (CH₂Cl₂:acetone = 19:1). Resultant product is recrystallized from hexane to give the aimed product [20a] (6.51g, 58%).

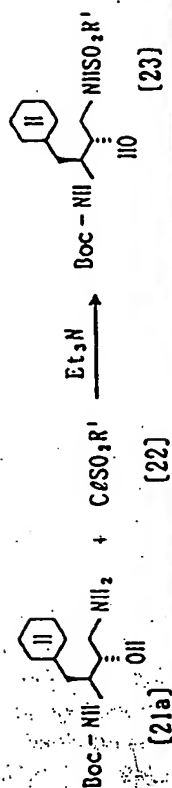
The product [20a] (3.56g, 12.6mmol) in anhydrous THF (50ml) is added dropwise to a suspension of LiAlH₄ (574mg, 1.2mol) in anhydrous THF (30ml) with stirring and ice-cooling over 30 minutes. The mixture is stirred at 0°C for an additional one hour. A small amount of ethyl acetate and ice water are added to the mixture to separate out inorganic materials. The insoluble materials are filtered, and the filtrate is concentrated *in vacuo* and then purified with silica gel chromatography (SiO₂: 120g, CH₂Cl₂:MeOH:NH₄OH = 80:20:2). The aimed compound [21a] (2.21g, 61%) is thus obtained.

To a solution of the compound [21a] (12.49g, 43.6mmol) in anhydrous dichloromethane (200ml) are added triethylamine (8.8g, 2.0eq) and morpholiniosulfonyl chloride (10.1g, 1.25eq) and the mixture is stirred at room temperature for 3 hours and concentrated *in vacuo*. The residue is dissolved in ethyl acetate, washed with water, dried, and evaporated to remove the solvent. The residue is purified with silica gel chromatography (SiO₂: 200g, CH₂Cl₂:MeOH:NH₄OH = 90:10:1). The aimed compound [23a] (18.16g, 95%) is thus obtained.

Preparation 65-74

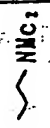


The compounds [23] listed in Table 6 are prepared in the manner as taught in Preparation 64.

Table 6

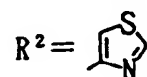
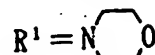
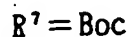
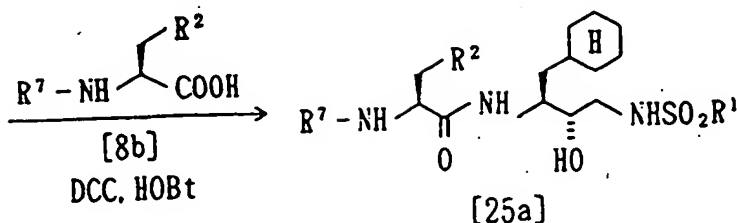
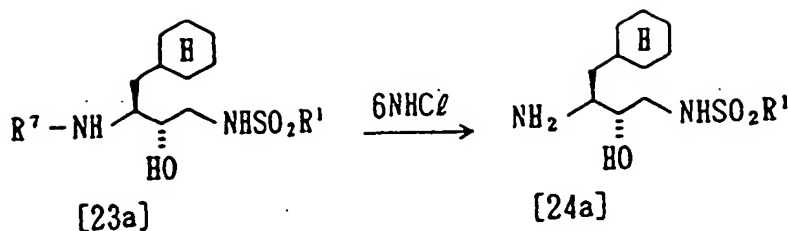


Compd. of prec. No.	R'	yield %	[23]	
			NMR (δ)	
65	-NMe ₂	97	0.80-1.90(13H, m), 1.45(9H, s), 2.80(6H, s), 3.08(2H, m), 3.72(2H, m), 4.63(1H, d, J=9.2Hz), 5.48(1H, m)	
66		90	0.70-1.80(13H, m), 1.37(9H, s), 2.49(1H, bs), 2.82(1H, dt, J=6.2, 13.5Hz), 3.12(1H, dt, J=7.13Hz), 3.68(2H, m), 4.60(1H, d, J=9.3Hz), 4.47(1H, dd, J=4.6Hz), 8.17(1H, m), 8.79(1H, bd), 9.08(1H, bs)	
67		93	0.70-1.85(13H, m), 1.39(9H, s), 2.90(1H, dt, J=6.1, 13.4Hz), 3.16(1H, dt, J=7.0, 12.9Hz), 2.60(1H, m), 3.67(2H, m), 4.57(1H, d, J=9.2), 5.89(1H, t, J=7Hz), 7.07(1H, dd, J=3.7, 5.0Hz), 7.60(2H, m), 6.24(1H, bt)	
68		99	0.65-1.80(13H, m), 1.32(9H, s), 2.89(2H, bt), 3.55(1H, m), 3.66(1H, m), 4.58(1H, d, J=9.2Hz), 6.80(1H, bt), 7.57(1H, dd, J=4.8, 4Hz), 7.66(1H, t, J=7.4Hz), 8.07(1H, dd, J=1.0, 8.2Hz), 8.29(1H, dd, J=1.6, 8.4Hz), 8.42(1H, dd, J=1.4, 7.4Hz), 9.05(1H, dd, J=1.6, 4.4Hz)	
69		97	0.70-1.80(13H, m), 1.37(9H, s), 2.58(1H, bd, J=5Hz), 2.80(1H, dt, J=6.2, 13.6Hz), 3.08(1H, dt, J=6.5, 13.6Hz), 3.65(2H, m), 4.56(1H, d, J=9.2Hz), 5.82(1H, bt, J=6Hz), 7.53(3H, m), 7.85(2H, m)	
70		72	0.75-1.87(13H, m), 1.45(9H, s), 1.98(1H, bs), 2.52(4H, m), 2.86(2H, m), 3.17(4H, m), 3.73(6H, m), 4.62(1H, d, J=9.4Hz), 5.63(1H, bt)	
71		36	0.75-1.88(13H, m), 1.45(9H, s), 2.03(2H, m), 2.49(6H, m), 3.13(4H, m), 3.72(6H, m), 4.65(1H, d, J=9.1Hz), 5.82(1H, bt)	

Table 6 (continued)

Compd. of Prep. No.	R'	yield %	[23]	
			NMR (δ)	
72		75	0.75-1.90(13H, m), 1.44(9H, s), 2.29(6H, s), 2.80(2H, m), 3.16(5H, m), 3.64(2H, m), 4.66(1H, d, J=9.6Hz)	
73		97	0.75-1.85(13H, m), 1.45(9H, s), 2.85(1H, bs), 2.96(3H, s), 3.14(2H, m), 3.72(2H, m), 4.65(1H, d, J=9.4Hz), 5.53(1H, bt)	
74		98	0.94(3H, t, J=7.2Hz), 0.80-1.95(17H, m), 1.45(9H, s), 2.76(1H, bs), 3.00(2H, m), 3.16(2H, m), 3.70(2H, m), 4.63(1H, d, J=9Hz), 5.44(1H, t, J=7Hz)	

Preparation 75



A mixture of the compound [23a] (18.16g, 41.6mmol), THF (150ml), and 6N HCl (150ml) is stirred at room temperature for 4 hours. The reaction mixture is made alkaline with Na_2CO_3 and saturated aqueous NaHCO_3 and extracted with a mixture of dichloromethane and methanol (9:1). The organic layer is dried and evaporated to dryness *in vacuo*. The residue is subjected to silica gel column chromatography (SiO_2 : 100g, CH_2Cl_2 :MeOH: NH_4OH = 80:20:2). The compound [24a] (14.0g, quantitative amount) is thus obtained.

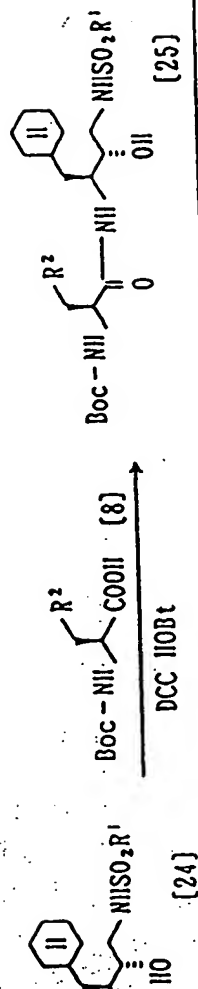
To a solution of the above compound [24a] (14.0g, 41.6mmol) in acetonitrile (200ml) are added 4-thiazolyl-L-alanine [8b] (12.09g, 1.1eq) and HOBt (7.04g, 1.25eq) with ice-cooling. To the mixture is added DCC (11.18g, 1.3eq) and the resulting mixture is stirred for one hour at 0 °C and one hour at room temperature. The reaction mixture is filtered after addition of ethyl acetate and the filtrate is concentrated *in vacuo*. The residue is subjected to silica gel column chromatography (SiO_2 : 600g, CH_2Cl_2 :MeOH: NH_4OH = 90:10:1) to give the product [25a] (24.5g, quantitative amount).

NMR(δ): 0.70-1.80(13H,m), 1.45(9H,s), 2.45(1H,bs), 2.98(2H,m), 3.18(4H,m), 3.30(2H,m), 3.75(5H,m), 4.02(1H,m), 4.46(1H,ddd,J=6.4Hz), 5.72(1H,bt,J=6.6Hz), 6.16(1H,d,J=6.4Hz), 6.36(1H,d,J=9.2Hz), 7.15(1H,d,J=1.8Hz), 8.82(1H,d,J=2Hz).

Preparation 76-86

Compounds [25] listed in Table 7 are prepared according to the procedure disclosed in Preparation 75.

Table 7



Compd. of Prep. No.	R ¹	R ²	Yield%	[25]	
				NMR (δ)	
76	-NMe ₂		90	0.70-1.80(13H, m), 1.45(9H, s), 2.79(6H, s), 2.95(2H, m), 3.29(2H, m), 3.73(1H, m), 4.01(1H, m), 4.48(1H, ddd, J=6.6Hz), 5.58(1H, bt), 6.15(1H, d, J=7Hz), 6.39(1H, d, J=10Hz), 7.15(1H, d, J=1.8Hz), 8.82(1H, d, J=2Hz)	
77			94	0.65-1.75(13H, m), 1.39(9H, s), 2.80(1H, dt, J=6.4, 13.6Hz), 3.00(1H, dt, J=6.6, 13.7Hz), 3.19(2H, d, J=5.6Hz), 3.70(1H, dt, J=2.3, 6.7Hz), 3.99(1H, m), 4.46(1H, ddd, J=6Hz), 6.05(1H, d, J=6.3Hz), 6.55(2H, m), 7.11(1H, d, J=1.8Hz), 7.45(1H, m), 7.80(1H, dd, J=4.24Hz), 8.20(1H, d, J=7Hz), 8.77(1H, d, J=2Hz), 9.08(1H, bs)	
78			99	0.65-2.00(13H, m), 1.43(9H, s), 2.81(1H, dt, J=6.3, 13.5Hz), 2.99(1H, dt, J=6.9, 13.5Hz), 3.24(2H, m), 3.66(1H, dd, J=2.4, 6.8Hz), 3.97(1H, m), 4.45(1H, ddd, J=6.5Hzx3), 6.02(1H, d, J=6.9Hz), 6.22(1H, bt), 6.39(1H, d, J=9.3Hz), 7.10(2H, m), 7.58(2H, m), 8.75(1H, d, J=1.8Hz)	
79			96	0.55(13H, m), 1.43(9H, s), 2.79(2H, m), 3.11(1H, dd, J=5.7, 14.7Hz), 3.22(1H, dd, J=5.4, 14.7Hz), 3.65(1H, m), 3.84(1H, m), 4.32(1H, ddd, J=6.6Hzx3), 6.09(1H, d, J=6Hz), 6.27(1H, d, J=9.4Hz), 6.78(1H, t, J=6.2Hz), 7.02(1H, d, J=1.8Hz), 7.56(1H, dd, J=4.3, 8.4Hz), 7.65(1H, t, J=7.4Hz), 8.06(1H, dd, J=1.4, 8.3Hz), 8.27(1H, dd, J=1.8, 8.4Hz), 8.41(1H, dd, J=1.4, 7.3Hz), 8.69(1H, d, J=1.9Hz), 9.04(1H, dd, J=1.7, 4.2Hz)	

Table 7 (continued)














Compd. of Prod. No.	R ¹	R ²	[25]	
			Yield%	NMR (δ)
80			99	0.67-2.00(13H, m), 1.43(9H, s), 2.74(1H, dt, J=6.9, 13.5Hz), 2.94(1H, dt, J=6.8, 13.5Hz), 3.18(1H, dd, J=6.3, 14Hz), 3.27(1H, dd, J=5.7, 14Hz), 3.62(1H, dt, J=2.6, 6.8Hz), 3.95(1H, m), 4.41(1H, ddd, J=6.6, 6Hzx3), 5.95(1H, bt), 6.05(1H, d, J=6.8), 6.29(1H, d, J=9.3Hz), 7.07(1H, d, J=1.9Hz), 7.52(3H, m), 7.86(2H, m), 8.73(1H, d, J=2.0Hz)
81			99	0.63-1.78(13H, m), 1.45(9H, s), 2.14(1H, bs), 2.52(4H, bt, J=4.6Hz), 2.86(2H, t, J=7Hz), 3.05(2H, bt, J=6Hz), 3.14-3.40(4H, m), 3.66(1H, m), 3.73(4H, m), 4.00(1H, m), 4.42(1H, ddd, J=6.2Hz), 5.80(1H, bt), 6.24(1H, d, J=6.6Hz), 6.31(1H, d, J=9.4Hz), 7.14(1H, d, J=2Hz), 8.81(1H, d, J=2Hz)
82			99	0.70-1.80(13H, m), 1.45(9H, s), 2.01(2H, m), 2.49(6H, m), 3.08(4H, m), 3.30(2H, m), 3.72(5H, m), 4.00(1H, m), 4.43(1H, ddd, J=6.6Hzx3), 5.88(1H, bt), 6.24(1H, d, J=6.6Hz), 6.52(1H, d, J=9.6Hz), 7.15(1H, d, J=1.8Hz), 8.82(1H, d, J=1.8Hz)
83			77	0.60-1.80(13H, m), 1.45(9H, s), 2.29(6H, s), 2.81(2H, t, J=6.2Hz), 3.04(2H, d, J=6.6Hz), 3.18(2H, m), 3.25(1H, dd, J=5.6, 14.6Hz), 3.34(1H, dd, J=5.4, 14.6Hz), 3.65(1H, dt, J=2.4, 6.2Hz), 3.98(1H, m), 4.45(1H, ddd, J=6.6Hz), 6.22(1H, d, J=6.6Hz), 6.40(1H, d, J=9.6Hz), 7.14(1H, d, J=2Hz), 8.81(1H, d, J=2Hz)
84	Me		75	0.70-1.80(13H, m), 1.45(9H, s), 2.96(6H, s), 3.03(2H, m), 3.30(2H, m), 3.70(1H, m), 4.02(1H, m), 4.46(1H, ddd, J=6.6Hzx3), 5.72(1H, bt), 6.21(1H, d, J=6.6Hz), 6.39(1H, d, J=9.6Hz), 7.15(1H, d, J=1.6Hz), 8.82(1H, d, J=1.8Hz)
85			83	0.95(3H, t, J=7.2Hz), 0.65-1.88(17H, m), 1.45(9H, s), 3.00(5H, m), 3.30(2H, m), 3.68(1H, dt, J=2.3, 6.6Hz), 4.01(1H, m), 4.45(1H, ddd, J=6.2Hzx3), 5.56(1H, bt), 6.18(1H, d, J=6.6Hz), 6.35(1H, d, J=9.6Hz), 7.14(1H, d, J=1.8Hz), 8.81(1H, d, J=2.0Hz)

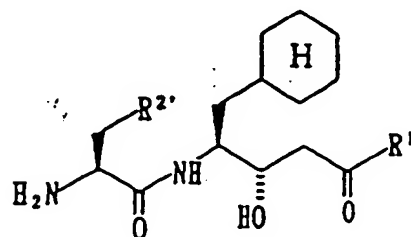
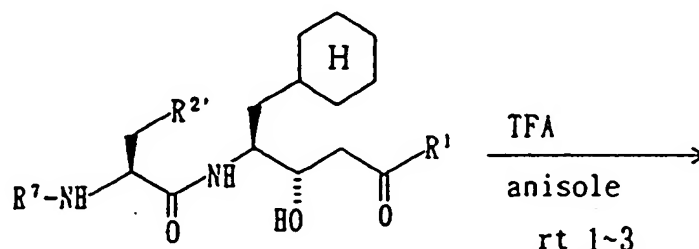
Table 7 (continued)

Compd. of Prep. No.	R ¹	R ²	Yield%	[25]	
				NMR (δ)	
86			99	0.65~1.75(13H, m), 1.43(9H, s), 2.64(3H, s), 2.74(1H, dt, J=6.3, 13.6Hz), 2.96(1H, dt, J=6.8, 13.4Hz), 3.10(2H, m), 3.61(1H, dt, J=3.6, 6Hz), 3.96(1H, m), 4.34(1H, ddd, J=5.8Hz), 5.90(1H, m), 6.00(1H, d, J=6.2Hz), 6.32(1H, d, J=9.2Hz), 6.82(1H, s), 7.52(3H, m), 7.86(2H, dd, J=1.6, 7.8Hz)	

Example 1

3-t-Butylsulfonyl-2(S)-phenylmethylpropionyl-His-1(S)-cyclohexylmethyl-2(S)-hydroxy-4-oxo-4-(4-pyridyl)butylamide [1a]

1) His(Ts)-1(S)-cyclohexylmethyl-2(S)-hydroxy-4-oxo-4-(4-pyridyl)butylamide [11a]



$R^1 = 4\text{-pyridyl}$

$R^2 = 4\text{-pyridyl}$

$R^{2'} = 4\text{-tosylimidazolyl}$

$R^7 : \text{Boc}$

Boc-His(Ts) 1(S)-cyclohexylmethyl-2(S), hydroxy-4-oxo-4-(4-pyridyl)butylamide [10a] (1.31g, 1.96mmol) prepared in Preparation 21 is dissolved in anisole (13ml). To the solution is added trifluoroacetic acid (13ml) with stirring and ice-cooling and the mixture is stirred at room temperature for one hour. After evaporation of the reaction mixture to dryness in vacuo, ice is added to the residue and the mixture is washed with ethyl ether. The aqueous layer neutralized with 3N NaOH and adjusted to pH8 by addition of powdered Na_2CO_3 is extracted with dichloromethane three times and finally extracted with a mixture of dichloromethane and methanol (10:1). The organic layer is washed with saturated aqueous sodium chloride, dried over MgSO_4 and evaporated to dryness in vacuo. The residue is purified with silica gel chromatography ($\text{CH}_2\text{Cl}_2:\text{MeOH} = 95:5$) to obtain the aimed crude product (850mg, 73%). Recrystallization of the crude product from ethyl acetate provides the title compound [11a] (750mg, 65%) as a needle crystal. m.p. 161-162°C

NMR(δ): 0.75-1.80(13H,m), 1.98(1H,br.s), 2.44(3H,s), 2.73(1H,dd,J=14.8,8.2Hz), 2.95-3.24(3H,m), 3.65(1H,dd,J=8.4,4.2Hz), 4.02(1H,m), 4.27(1H,m), 7.12(1H,d,J=1.2Hz), 7.36(2H,d,J=7.8Hz), 7.53(1H,d,J=10Hz), 7.70(2H,m), 7.81(2H,d,J=8.4Hz), 7.92(1H,d,J=1.4Hz), 8.79(2H,m)

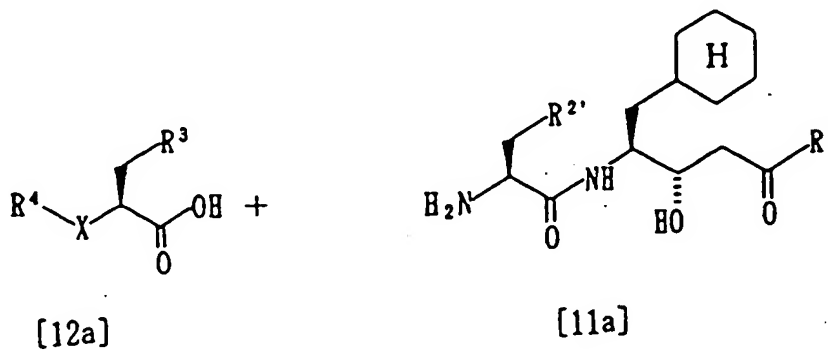
IR $\nu_{\text{max}}(\text{CHCl}_3)\text{cm}^{-1}$: 3680, 3340, 1690, 1654, 1602, 1593, 1515, 1475, 1450

Elemental analysis(as $\text{C}_{29}\text{H}_{39}\text{N}_5\text{O}_6\text{S}$)

Calcd.: C:59.01; H:6.75; N:11.87; S:5.43

Found : C:59.12; H:6.69; N:11.68; S:5.21

2) 3-t-Butylsulfonyl-2(S)-phenylmethylpropionyl-His(Ts)-1(S)-cyclohexylmethyl-2(S)-hydroxy-4-oxo-4-(4-pyridyl)butylamide [13a]



To a solution of the protected compound [13a] (740mg, 0.89mmol) obtained in the above step 2) in DMF (4ml) is added pyridinium hydrochloride (1030mg, 8.87mmol, 10.0eq) and the mixture is stirred at room temperature for two hours. The reaction mixture is adjusted to pH 7 - 8 by addition of ice and 4% aqueous NaHCO₃ and extracted three times with dichloromethane. The organic layer is washed with saturated aqueous sodium chloride, dried over MgSO₄, and concentrated to dryness in vacuo. The residue is purified with silica gel chromatography (CH₂Cl₂:MeOH:concNH₄OH = 950:50:1) to obtain the title compound [1a] (543mg, 90%). Trituration of the residue with diisopropyl ether gives colorless powders.

NMR δ : 0.67-1.83(13H,m), 1.33(9H,s), 2.86(1H,d,J=13.5,8.4Hz), 2.97(1H,dd,J=13.0,9.8Hz), 3.10(5H,m), 3.26(1H,m), 3.56(1H,dd,J=13.0,9.8Hz), 4.02(1H,m), 4.20(1H,m), 4.56(1H,ddd,J=6.3,6.3,6.3Hz), 6.44 (1H, d, J=10Hz), 6.90(1H,s), 7.24(4H,m), 7.48(1H,s), 7.46(1H,bs), 7.70(2H,m), 8.78(2H,m)
[α]_D = -22.5° (C=1.0; MeOH; 23°C) IR v_{max}(CHCl₃)cm⁻¹: 3460, 3360(br), 1662(1690sh), 1603, 1496, 1450, 1410, 1115

Elemental analysis (as C₃₈H₄₉N₅O₆S.3/4H₂O)

Calcd.: C:62.36; H:7.34; N:10.10; S:4.62

Found : C:62.42; H:7.33; N:10.21; S:4.49

Examples 2-52

The same procedure as disclosed in the steps 1) and 2) in Example 1 is repeated using, as the starting material, the compounds [10] prepared in foregoing Preparations 21-58, and the compounds [11] and [13] listed in Tables 8 (compound [11]) and 9 (compound [13]) are obtained. The compounds [13] (for example, compound [13] of No. 23) wherein R¹ or R² is not protected correspond to the compounds (I) of the invention. Where the substituent R² is protected, the compounds [13] are deprotected according to the procedure as disclosed in Step 3) in Example 1 to obtain the final products (I), which are listed in the following Table 10.

Table 8







Comp. of Ex. No.	R ¹	R ²	(11)			IR ν _{max} cm ⁻¹ or NMR (δ)	Elemental analysis	
			Yield%	[α] _D ²⁰ (C=1.0, CHCl ₃) (°C)	a.p. (°C)		Calcd.	Found
2	phenyl		81	-49.1 (23.5)	126-127	3560, 3360, 1666, 1598, 1580, 1511 1450, 1382, 1172, 1075	C: 63.58 H: 6.76 N: 9.89 S: 5.66	C: 63.34 H: 6.67 N: 9.84 S: 5.67
3	o-fluorophenyl		67			3600, 3460(br), 1670, 1610, 1598, 1510, 1480, 1450, 1383, 1170, 1075	oil	
4	m-methoxyphenyl		55			0.70-1.83(13H, m), 2.05(3H, bs), 2.44(3H, s), 2.72(1H, dd, J=8, 16Hz), 2.98(1H, dd, J=10, 18Hz), 3.07(1H, dd, J=6, 16Hz), 3.18(1H, dd, J=3, 18Hz), 4.66(1H, dd, J=4, 8Hz), 3.85(3H, s), 4.02(1H, m), 4.24(1H, m), 7.17(1H, s), 7.17(1H, m), 7.30-7.60(3H, m), 7.35(2H, d, J=8, 0Hz), 7.80(2H, d, J=8, 4Hz), 7.91(1H, d, J=1, 4Hz)	oil	
5	p-methylphenyl		43			3360, 1668, 1608, 1570, 1510, 1450, 1385, 1172, 1092, 1075	oil	
6	2,4-difluorophenyl		71	-43.9 (23.5)		3580, 3360, 1665, 1612, 1595, 1510, 1498, 1475, 1383, 1172, 1075, 970	C: 59.78 H: 6.02 N: 9.30 S: 5.32 F: 6.31	C: 59.53 H: 6.04 N: 9.42 S: 5.56 F: 6.38
7	1-naphthyl		49			3400(br), 1665, 1599, 1575, 1510, 1450, 1386, 1190, 1174, 1094, 1080, 909	oil	

Table 8 (continued)

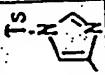


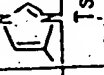
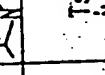

Compd. of Ex. No.	R ¹	R ²	[11]		mp(°C)	IR ν _{max} cm ⁻¹ or NMR(δ)
			Yield%	[α] _D ²⁰ (C=1.0, CHCl ₃)(°)		
8	3-thienyl		70			3680, 3360(br), 3120, 1665, 1598, 1510, 1475, 1450, 1382, 1172, 1076
9	2-thiazolyl		52			0. 73-1. 83(13H, m), 2. 33(2H, bs), 2. 44(3H, s), 2. 76(1H, dd, J=7, 15Hz), 3. 07(1H, dd, J=3, 6, 14, 6Hz), 3. 18-3. 38(2H, m), 3. 68(1H, m), 4. 02(1H, m), 4. 25(1H, m), 7. 14(1H, s), 7. 35(2H, d, J=8, 0Hz), 7. 51(1H, d, J=9, 0Hz), 7. 69(1H, d, J=3, 0Hz), 7. 80(2H, d, J=8, 4Hz), 7. 95(1H, d, J=1, 2Hz), 8. 01(1H, d, J=3, 0Hz)
10	m-fluorophenyl		66			3360, 1670, 1590, 1510, 1445, 1382, 1170, 1090, 1075
11	p-fluorophenyl		57	-45.8 (24.0)	128-130	3360, 3500(br), 1665, 1600, 1508, 1475, 1450 1095, 1075
12	2,6-difluoro- phenyl		53	-23.9 (24.0)		3368, 1698, 1665, 1624, 1598, 1512, 1420, 1385, 1279, 1190, 1174, 1094, 1077, 1018
13	o-acethoxyphenyl		25			0. 70-1. 85(13H, m), 2. 20(3H, bs), 2. 44(3H, s), 2. 74(1H, dd, J=8, 15Hz), 2. 95(1H, dd, J=10, 17Hz), 3. 10(1H, dd, J=15, 5Hz), 3. 26(1H, dd, J=17, 3Hz), 3. 69(1H, dd, J=5, 10Hz), 3. 88(3H, s), 3. 99(1H, m), 4. 18(1H, m), 6. 98(2H, m), 7. 12(1H, s), 7. 35(2H, d, J=8Hz), 7. 50(1H, m), 7. 72(1H, dd, J=7, 5, 2Hz), 7. 81(2H, d, J=8Hz), 7. 92(1H, s)

Table 8 (continued)

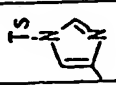
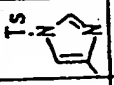


Compd. of Ex. No.	R ¹	R ²	[11]		IR ν_{\max} cm ⁻¹ or NMR (δ)
			Yield%	mp (°C)	
14	o-chlorophenyl		75		0.70-1.82 (13H, m), 2.40 (3H, s), 2.44 (3H, s), 2.72 (1H, dd, J=8, 15, 8Hz), 3.00 (1H, dd, J=17, 5, 10Hz), 3.07 (1H, dd, J=15, 5Hz), 3.17 (1H, dd, J=17, 5, 4Hz), 3.65 (1H, dd, J=10, 5Hz), 3.98 (1H, m), 4.23 (1H, m), 7.11 (1H, s), 7.25-7.58 (6H, m), 7.81 (2H, d, J=8, 4Hz), 7.91 (1H, d, J=1, 4Hz)
15	m-cyanophenyl		79		3360, 2236, 1666, 1514, 1498, 1450, 1386, 1189, 1174, 1094, 1078, 909
16	o-ethyl-sulfonyl-aminophenyl		38		3368, 1657, 1607, 1578, 1496, 1452, 1386, 1340, 1189, 1173, 1094, 1078, 968, 909
17	p-trifluoro-methylphenyl		53	113-115	3360, 1670, 1600, 1510, 1450, 1410, 1385, 1325, 1180, 1135, 1065

Table 8 (continued)






Compd. of Ex. No.	R ¹	R ²	(II)	
			Yield%	NMR (δ)
18	m-morpholino- carbonyloxy- phenyl		81	0. 73-2. 20(13H, m), 2. 44(3H, s), 2. 75(1H, dd, J=14. 8, 8. 6Hz), 2. 93-3. 24(3H, m), 3. 5-3. 82(8H, m), 4. 02(1H, m), 4. 23(1H, m), 7. 13(1H, d, J=1. 0Hz), 7. 35(2H, d, J=8. 0Hz), 7. 35(1H, m), 7. 47(1H, t, J=7. 5Hz), 7. 60(1H, d, J=10Hz), 7. 67(1H, m), 7. 81(2H, d, J=8. 4Hz), 7. 81(1H, m), 7. 91(1H, d, J=1. 4Hz)
19	m-morpholino- carbonylophenyl		41	0. 70-1. 85(13H, m), 2. 28(3H, bs), 2. 44(3H, s), 2. 75 (1H, dd, J=8. 6, 14. 8Hz), 2. 95-3. 27(3H, m), 3. 30-3. 94(8H, m), 3. 68 (1H, dd, J=8. 4, 2Hz), 4. 02(1H, m), 4. 26(1H, m), 7. 14(1H, d, J=1. 4Hz), 7. 36(2H, d, J=8Hz), 7. 47-7. 69(2H, m), 7. 81(2H, d, J=8. 4Hz), 7. 93(1H, d, J=1. 2Hz), 7. 98(1H, d, J=1. 6Hz), 8. 00(1H, m)
20	3,4-ethylene- dioxyphenyl		74	0. 70-2. 15(13H, m), 2. 44(3H, s), 2. 73(1H, dd, J=14. 4, 8. 4Hz), 2. 91(1H, dd, J=17. 8, 9. 6Hz), 3. 09(1H, dd, J=14. 6, 4. 2Hz), 3. 13(1H, dd, J=18. 4Hz), 3. 67(1H, dd, J=8. 6, 3. 8Hz), 4. 00(1H, m), 4. 20(1H, m), 6. 05(2H, s), 6. 84(1H, d, J=8. 2Hz), 7. 12(1H, d, J=1. 0Hz), 7. 36(2H, d, J=8. 0Hz), 7. 40(1H, d, J=1. 6Hz), 7. 53(1H, dd, J=8. 2, 1. 6Hz), 7. 81(2H, d, J=8. 2Hz), 7. 92(1H, d, J=1. 4Hz)
21	cyclohexyl		68	0. 70-1. 89(23H, m), 2. 13(3H, bs), 2. 33(1H, m), 2. 45(1H, m), 2. 47(1H, dd, J=17. 6, 9. 4Hz), 2. 66(1H, dd, J=15. 2, 6Hz), 2. 71(1H, dd, J=14. 9, 4Hz), 3. 07(1H, dd, J=14. 8, 3. 6Hz), 7. 12(1H, d, J=1. 2Hz), 7. 37(2H, d, J=8. 4Hz), 7. 48(1H, d, J=10Hz), 7. 82(2H, d, J=8. 4Hz), 7. 94(1H, d, J=1. 4Hz)
22	p-ethoxyphenyl		51	0. 76-2. 20(13H, m), 2. 44(3H, s), 2. 72(1H, dd, J=8. 6, 15Hz), 2. 93(1H, dd, J=9. 6, 17. 6Hz), 3. 94(1H, dd, J=3. 6, 15Hz), 3. 17(1H, dd, J=2. 4, 17. 6Hz), 3. 67(1H, dd, J=4. 8, 6Hz), 3. 88(3H, s), 4. 02(1H, m), 4. 23(1H, m), 6. 93(2H, d, J=9Hz), 7. 27(1H, s), 7. 36(2H, d, J=8. 2Hz), 7. 52(1H, d, J=9. 6Hz), 7. 81(1H, d, J=8. 4Hz), 7. 91(2H, d, J=9Hz), 7. 92(1H, d, J=1. 8Hz)

Table 8 (continued)

Compd. of Ex. No.	R ¹	R ²	[11]	
			Yield%	NMR (δ)
23	phenyl		70	0.7-2.05(13H, m), 2.96(1H, dd, J=18, 9, 4Hz), 3.15(1H, dd, J=14, 2, 7, 8Hz), 3.21(1H, dd, J=18, 2, 6Hz), 3.36(1H, dd, J=14, 2, 4, 2Hz), 3.80(1H, dd, J=7, 8, 4, 4Hz), 4.04(1H, m), 4.24(1H, m), 7.11(1H, d, J=1, 6Hz), 7.41-7.63(3H, m), 7.94(2H, m), 8.75(1H, d, J=1, 8Hz) (mp. 106-107°C)
24	4-pyridyl		70	0.70-1.85(13H, m), 2.04(3H, m), 3.02(1H, dd, J=18, 8, 6Hz), 3.10- 3.26(2H, m), 3.36(1H, dd, J=14, 4, 4, 2Hz), 3.82(1H, dd, J=7, 6, 4, 4Hz), 4.02(1H, m), 4.26(1H, m), 7.13(1H, d, J=1, 6Hz), 7.59(1H, d, J=10Hz), 7.71(2H, dd, J=4, 6, 1, 6Hz), 8.76(1H, d, J=2Hz), 8.82(2H, dd, J=4, 6, 1, 6Hz) (mp. 118-120°C)
25	3-thienyl		72	0.70-1.87(13H, m), 2.28(3H, bs), 2.89(1H, dd, J=17, 6, 9, 4Hz), 3.10(1H, dd, J=17, 6, 2, 7Hz), 3.14(1H, dd, J=14, 3, 7, 8Hz), 3.35(1H, dd, J=14, 3, 4, 1Hz), 3.78(1H, dd, J=7, 8, 4, 3Hz), 4.00(1H, m), 4.20(1H, m), 7.12(1H, d, J=2, 0Hz), 7.31(1H, dd, J=5, 1, 2, 9Hz), 7.52(1H, dd, J=5, 1, 1, 2Hz), 7.57(1H, s), 8.08(1H, dd, J=2, 9, 1, 2Hz), 8.75(1H, dd, J=2Hz)
26	cyclohexyl		69 mp. 90-93	0.70-1.90(24H, m), 1.98(3H, bs), 2.32(1H, m), 2.45(1H, dd, J=9, 8, 1, 8Hz), 2.66(1H, dd, J=18, 2, 8Hz), 3.15(1H, dd, J=14, 7, 4Hz), 3.35(1H, dd, J=14, 4, 3, 8Hz), 3.80(1H, dd, J=7, 4, 4, 2Hz), 3.91(1H, m), 4.02(1H, m), 7.13(1H, d, J=1, 6Hz), 7.50(1H, d, J=9, 8Hz), 8.78(1H, d, J=1, 8Hz)

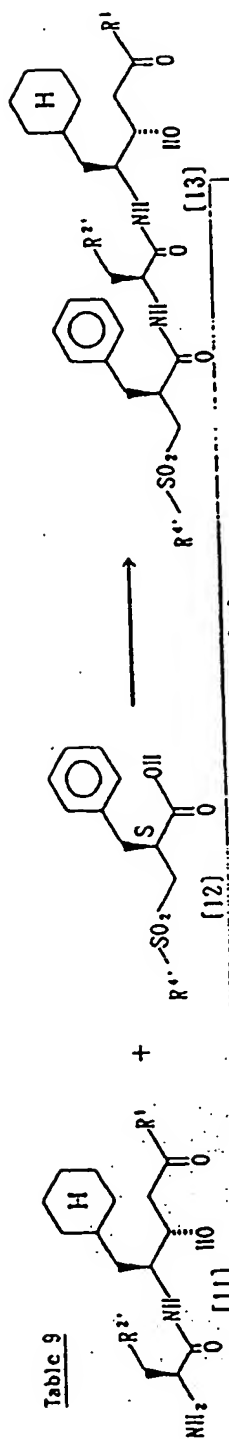
Table 8 (continued)

Compd. of Ex. No.	R ¹	R ²	(II)		
			Yield%	[α] _D ²⁰ (C=1, CHCl ₃) (Temp. °C)	NMR(δ)
27	m-2-(N-morpholino)-ethoxyphenyl		76	-46.8 (23.5)	0.70-2.10(13H, m), 2.59(4H, t, J=4.7 Hz), 2.82(2H, t, J=5.7 Hz), 2.94(1H, dd, J=9.5, 17.9 Hz), 3.15(1H, dd, J=7.6, 14.6 Hz), 3.18(1H, dd, J=2.5, 17.9 Hz), 3.34(1H, dd, J=4.1, 14.6 Hz), 3.74(4H, t, J=9.3 Hz), 3.74(1H, m), 4.02(1H, m), 4.15(2H, t, J=8.7 Hz), 4.22(1H, m), 7.11(1H, br, s), 7.15(1H, d, J=2.7), 7.36(1H, t, J=8.2), 7.45-7.62(2H, m), 8.75(1H, d, J=1.1 Hz)
28	m-(N-forayl)-methylamino-phenyl		73		0.77-1.85(13H, m), 2.40(2H, m), 3.02(1H, dd, J=9.0, 17.9 Hz), 3.17(3H, m), 3.34(3H, s), 3.35(1H, m), 3.60(1H, dd, J=4.3, 7.8 Hz), 4.03(1H, m), 4.25(1H, dt, J=2.1, 8.2 Hz), 7.13(1H, d, J=2.0 Hz), 7.38(1H, ddof d, J=8.0, 1.2, 2.3 Hz), 7.52(1H, t, J=7.52 Hz), 7.59(1H, d, J=9.7 Hz), 7.79(2H, m), 8.52(1H, s), 8.75(1H, d, J=2.0)
29	N-methyl-3-pyrrolyl		69	-57.6 (24)	0.70-2.00(13H, m), 2.65(1H, dd, J=9.8, 16.8 Hz), 2.94(1H, dd, J=2.4, 17.1 Hz), 3.13(1H, dd, J=7.6, 14.4 Hz), 3.35(1H, dd, J=4.2, 14.6 Hz), 3.69(3H, s), 3.77(1H, dd, J=4.2, 8.4 Hz), 3.99(1H, m), 4.13(1H, dt, J=9.6, 2.1 Hz), 6.56(1H, s), 6.57(1H, s), 7.11(1H, d, J=1.8 Hz), 7.27(1H, s), 7.53(1H, d, J=9.6 Hz), 8.76(1H, d, J=2.1 Hz)
30	N-morpholino-methyl		52		
31	N-piperidino-methyl		36		
32	4-pyridyl		76		0.76-1.90(13H, m), 2.35(3H, bs), 2.67(3H, s), 3.14(4H, m), 3.80(1H, dd, J=4.2, 7.8 Hz), 4.03(1H, m), 4.25(1H, m), 6.88(1H, s), 7.58(1H, d, J=9.6 Hz), 7.71(2H, m), 8.81(2H, m)
33	phenyl		70		0.78-1.75(13H, m), 3.11(4H, m), 4.07(2H, m), 4.23(1H, m), 6.70(1H, s), 7.41(2H, dd, J=7.8, 15.3 Hz), 7.57(1H, t, J=7.1 Hz), 7.88(2H, d, J=7.2 Hz), 8.49(1H, s)

Table 8 (continued)

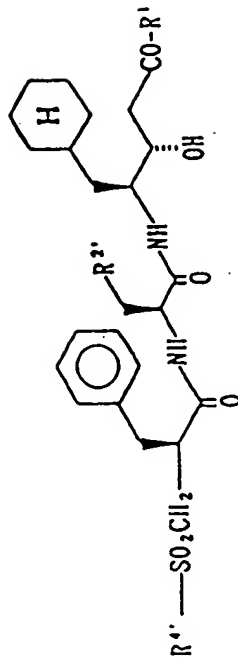
Compd. of Ex. No.	R ¹	R ²	[11]	
			Yield% (C=I, CHCl ₃)	[α] _D ²⁰ (Temp. °C)
34	4-pyridyl	-CONH ₂	17	
35	4-pyridyl	-SMe	88	
				0.70-1.80(13H, m), 2.13(3H, s), 2.73(1H, dd, J=8.2, 13.8Hz), 3.02(1H, dd, J=4.13, 6.1Hz), 3.09(1H, dd, J=8.6, 15.2Hz), 3.20(1H, dd, J=3.6, 18.4Hz), 3.58(1H, dd, J=4.8, 4.1Hz), 4.06(1H, m), 4.28(1H, m), 7.58(1H, d, J=10.1Hz), 7.72(2H, m), 8.81(2H, m)

Table 9



Compd. of Ex. No.	R ¹	R ²	R ³	IR ν _{max} (cm ⁻¹) or NMR(δ)	
				Yield %	
2	phenyl		tert-butyl	86	0.70-1.82(13H, m), 1.32(9H, s), 2.44(3H, s), 2.75-3.14(7H, m), 3.21(1H, m), 3.48(1H, dd, J=13.9Hz), 3.98(1H, m), 4.20(1H, m), 4.56(1H, ddd, J=6.6, 6.6Hz), 6.44(1H, d, J=9.4Hz), 7.13-7.30(6H, m), 7.40-7.63(3H, m), 7.34(2H, d, J=8.4Hz), 7.80(2H, d, J=8.4Hz), 7.89(1H, d, J=1.2Hz), 7.96(2H, d, J=7.8Hz)
3	o-fluorophenyl		tert-butyl	73	0.70-1.80(13H, m), 1.34(9H, s), 2.15(2H, bs), 2.70-3.15(7H, m), 3.19(1H, m), 3.51(1H, dd, J=9.6, 13.4Hz), 3.97(1H, m), 4.15(1H, m), 4.58(1H, ddd, J=6.2, 6.2, 2.2Hz), 6.43(1H, d, J=9.0Hz), 7.18(1H, s), 7.05-7.41(7H, m), 7.34(2H, d, J=8.2Hz), 7.54(1H, m), 7.82(2H, d, J=8.6Hz), 7.87(1H, ddd, J=7.7, 1.9Hz), 7.96(1H, d, J=1.7Hz)
4	o-acethoxyphenyl		tert-butyl	93	0.70-1.82(13H, m), 1.33(9H, s), 2.43(3H, s), 2.77-3.13(7H, m), 3.20(1H, m), 3.49(1H, dd, J=9.16Hz), 3.86(3H, s), 3.48(1H, m), 4.18(1H, m), 4.56(1H, ddd, J=7.7, 7.7Hz), 6.40(1H, d, J=9.0Hz), 7.21(1H, s), 7.09-7.40(7H, m), 7.52(2H, m), 7.80(2H, d, J=8.4Hz), 7.87(1H, d, J=1.4Hz)
5	p-ethylphenyl		tert-butyl	79	3400, 3260, 3140, 1665, 1625, 1605, 1498, 1450, 1370, 1172, 1115, 1030, 1010
6	2,4-difluoro-phenyl		tert-butyl	76	3400(br), 1655, 1600, 1599(ah), 1500, 1475, 1175, 970, 855
7	1-naphthyl		tert-butyl	41	3696, 3416, 1667, 1598, 1509, 1477, 1450, 1385, 1292, 1175, 1117, 1094, 1080

Table 9 (continued)



Compd. of Ex. No.	R'	R''	R'''	[13]	
				Yield %	$\nu_{max}(cm^{-1})$ or NMR(δ)
8	3-thienyl	tert-butyl		74	3410, 3360(sh), 1665, 1598, 1510, 1385, 1173, 1116, 1093, 1078
9	2-thiazolyl	tert-butyl		81	0, 7-1.82(13H, m), 1.35(9H, s), 2.37(1H, bs), 2.43(3H, s), 2.78-3.28(8H, m), 3.53(1H, dd, J=9.0, 13.0 Hz), 3.97(1H, m), 4.18(1H, m), 4.58(1H, ddd, 6.4, 6.4, 6.4 Hz), 6.35(1H, d, J=9.0 Hz), 7.05(1H, d, J=6.4 Hz), 7.20(1H, s), 7.13-7.40(5H, m), 7.33(2H, d, J=8.4 Hz), 7.68(1H, d, J=3.2 Hz), 7.80(2H, d, J=8.4 Hz), 8.00(1H, d, J=1.2 Hz), 8.01(2H, d, J=3.1 Hz)
10	m-fluorophenyl	tert-butyl		83	3460, 3360, 3280, 3160, 1665, 1625, 1590, 1500, 1450, 1115, 1032, 1010

Table 9 (continued)








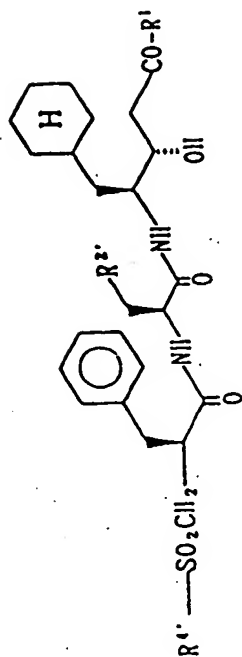
Compd. of Ex. No.	R ¹	R ⁴	R ²	Yield %	[13]	
					I R or NMR	
11	p-fluorophenyl	tert-butyl		71	3410, 3280, 3160, 1665, 1625, 1600, 1509, 1450, 1155, 1115, 1030, 1010	
12	2,6-difluoro-phenyl	tert-butyl		97	3420, 1660, 1624, 1599, 1499, 1467, 1459, 1385, 1292, 1189, 1175, 1118, 1093, 1085, 1018	
13	o-acetoxyphenyl	tert-butyl		79	0.70-1.80(13H, m), 1.33(9H, s), 2.42(3H, s), 2.78-3.25(7H, m), 3.50(1H, dd, J=18, 13Hz), 3.88(3H, s), 3.95(1H, m), 4.10(1H, m), 4.48(1H, ddd, J=6, 5, 6, 5Hz), 6.48(1H, d, J=9Hz), 6.97(2H, m), 7.06-7.40(9H, m), 7.49(1H, m), 7.73(1H, dd, J=9, 2Hz), 7.79(2H, d, J=8Hz), 7.84(1H, s)	
14	o-chlorophenyl	tert-butyl		81	0.70-1.80(13H, m), 1.31(9H, s), 2.40(3H, s), 2.80-3.22(7H, m), 3.50(1H, dd, J=15, 7, 5Hz), 3.93(1H, m), 4.09(1H, m), 4.51(1H, ddd, J=6, 4, 6, 4, 4Hz), 6.29(1H, d, J=10Hz), 7.03-7.41(8H, m), 7.53(1H, m), 7.78(2H, d, J=8, 4Hz), 7.78(1H, d, J=1, 4Hz)	
15	m-cyanophenyl	tert-butyl		84	3408, 2236, 1668, 1599, 1508, 1478, 1450, 1368, 1291, 1190, 1175, 1117, 1079, 908	
16	o-ethyl-sulfonyl-aminophenyl	tert-butyl			3420, 1666, 1607, 1578, 1499, 1452, 1387, 1340, 1290, 1174, 1155, 1117, 1079, 968, 909	
17	p-trifluoroacetylphenyl	tert-butyl		85	3400-3200, 3140, 1665, 1625, 1600, 1510, 1450, 1410, 1325, 1175, 1135, 1115, 1065	

Table 9 (continued)



Compd. of Ex. No.	R'	R''	R'''	Yield %	[13]	
					NMR (δ)	
18	m-morpholino- carbonyloxyphenyl	tert-butyl	Ts	79	0: 70-1. 82(13H, m), 1. 90(1H, bs), 1. 32(9H, s), 2. 43(3H, s), 2. 74-3. 30(7H, m), 3. 49(1H, dd, J=14, 10Hz), 3. 52-3. 82(8H, m), 3. 96(1H, m), 4. 16(1H, m), 4. 54(1H, ddd, J=6, 2, 6, 2Hz), 6. 40(1H, d, J=9, 4Hz), 7. 10-7. 40(7H, m), 7. 33(2H, d, J=8, 2Hz), 7. 48(1H, t, J=7, 5Hz), 7. 70(1H, m), 7. 80(2H, d, J=8, 4Hz), 7. 80(1H, m), 7. 89(1H, d, J=1, 2Hz)	
19	m-morpholino- carbonylphenyl	tert-butyl	Ts	85	0: 7-1. 82(13H, m), 1. 32(9H, s), 2. 43(3H, s), 2. 70-3. 30(8H, m), 3. 48(1H, dd, J=9, 4, 12, 8Hz), 3. 30-3. 90(8H, m), 3. 99(1H, m), 4. 20(1H, m), 4. 53(1H, ddd, J=6, 2, 6, 2, 6, 2Hz), 6. 42(1H, d, J=9, 4Hz), 7. 08-7. 31(6H, m), 7. 34(2H, d, J=8, 2Hz), 7. 45-7. 68(2H, m), 7. 81(2H, d, J=8, 4Hz), 7. 91(1H, d, J=1, 4Hz), 8. 05(2H, m)	
20	3',4'-methylene- dioxypheyl	tert-butyl	Ts	89	0: 7-1. 8(13H, m), 1. 9(1H, bs), 1. 33(9H, s), 2. 44(3H, s), 2. 74-3. 30(7H, m), 3. 50(1H, dd, J=9, 2, 13Hz), 3. 96(1H, m), 4. 16(1H, m), 4. 56(1H, ddd, J=6, 4, 6, 4, 6, 4Hz), 6. 06(2H, s), 6. 40(1H, d, J=9, 2Hz), 6. 85(1H, d, J=8, 2Hz), 7. 25(6H, m), 7. 34(1H, d, J=8, 2Hz), 7. 45(1H, d, J=1, 6Hz), 7. 56(1H, dd, J=8, 2, 1, 6Hz), 7. 80(2H, d, J=8, 4Hz), 7. 85(1H, d, J=1, 4Hz)	

Table 9 (continued)

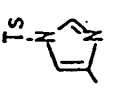

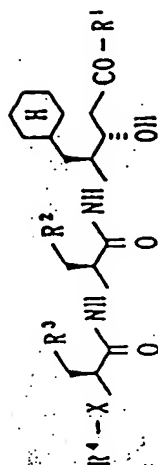
Compd. of Ex. No.	R ¹	R ⁴	R ²	Yield %	[13]	
					NMR (δ)	
21	cyclohexyl	tert-butyl		86	0. 70-1. 90(23H, m), 1. 34(9H, s), 2. 33(1H, m), 2. 44(3H, s), 2. 78-3. 27(8H, m), 3. 50(1H, dd, J=12. 0, 8. 4Hz), 3. 86(1H, m), 3. 96(1H, m), 4. 54(1H, ddd, J=6. 2, 6. 2, 6. 2Hz), 6. 35(1H, d, J=10Hz), 7. 13-7. 34(6H, m), 7. 37(2H, d, J=8. 4Hz), 7. 82(2H, d, J=8. 4Hz), 7. 90(1H, d, J=1. 4Hz).	
22	p-methoxyphenyl	tert-butyl		87	0. 72-1. 80(13H, m), 0. 90(1H, bs), 1. 33(9H, s), 2. 43(3H, s), 2. 78-3. 12(7H, m), 3. 20(1H, m), 3. 49(1H, dd, J=9. 4, 13. 2Hz), 3. 88(3H, s), 3. 97(1H, dd, J=9. 4, 13. 2Hz), 3. 88(3H, s), 3. 97(1H, ddd, J=7. 4, 7. 4, 7. 4Hz), 4. 17(1H, m), 4. 57(1H, ddd, J=6. 6, 6. 6Hz), 6. 42(1H, d, J=9. 4Hz), 6. 94(2H, d, J=9Hz), 7. 16-7. 26(6H, m), 7. 22(1H, s), 7. 34(2H, d, J=8Hz), 7. 79(2H, d, J=8. 4Hz), 7. 84(1H, d, J=1. 2Hz), 7. 94(2H, d, J=9Hz).	

Table 9 (continued)



Compd. of Ex. No.	R¹	R²	R³	X	R⁴	[13] ([1A])	
						Yield%	NMR (δ)
23	phenyl			CH₂	+SO₂	86	0.7-1.88(13H, m), 1.32(9H, s), 2.83-3.55(9H, m), 3.96(1H, m), 4.15(1H, m), 4.74(1H, ddd, J=5.6Hzx3), 6.32(1H, d, J=9.8Hz), 7.13(1H, d, J=2Hz), 7.25(5H, m), 7.47(2H, t, J=7.8Hz), 7.57(1H, d, J=7.8Hz), 7.96(2H, d, J=7.2Hz), 8.65(1H, d, J=2Hz)
24	4-pyridyl			CH₂	+SO₂	86	0.60-1.80(13H, m), 1.33(9H, s), 1.88(2H, bs), 2.86-3.50(9H, m), 3.99(1H, m), 4.15(1H, m), 4.66(1H, ddd, J=6Hzx3), 6.33(1H, d, J=9.2Hz), 7.16(1H, d, J=0.6Hz), 7.28(5H, m), 7.65(1H, d, J=6Hz), 7.82(2H, bs), 8.69(1H, d, J=0.6Hz), 8.82(2H, bs)
25	3-thienyl			CH₂	+SO₂	79	0.65-1.82(13H, m), 1.33(9H, s), 2.82-3.55(9H, m), 3.95(1H, m), 4.15(1H, m), 4.72(1H, ddd, J=6Hzx3), 6.33(1H, d, J=9.4Hz), 7.14(1H, d, J=2Hz), 7.17-7.38(6H, m), 7.55(1H, dd, J=5.2, 1.4Hz), 7.55(1H, s), 8.20(1H, dd, J=1.2, 2.8Hz), 8.67(1H, d, J=2Hz)
26	cyclohexyl			CH₂	+SO₂	87	0.7-2.00(24H, m), 1.35(9H, s), 2.32(1H, m), 2.88-3.58(9H, m), 3.83(1H, m), 3.92(1H, m), 4.70(1H, ddd, J=5.2Hzx3), 6.30(1H, d, J=9.8Hz), 7.16(1H, d, J=2Hz), 7.27(5H, m), 7.50(1H, d, J=5.8Hz), 8.71(1H, d, J=2.2Hz)
27	m-2-(N-morpholinoethoxy)phenyl			CH₂	+SO₂	91	0.6-1.8(13H, m), 2.60(4H, m), 2.84(2H, t, J=1.2Hz), 2.89-3.62(9H, m), 3.75(4H, m), 3.95(1H, ddd, J=7.8Hzx3), 4.18(2H, t, J=5.4Hz), 4.18(1H, m), 4.73(1H, ddd, J=5.2Hzx3), 6.32(1H, d, J=9.4Hz), 7.14(1H, d, J=2Hz), 7.14(1H, m), 7.22-7.56(3H, m), 8.67(1H, d, J=2Hz)

Table 9 (continued)



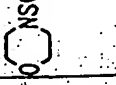




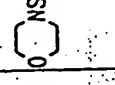


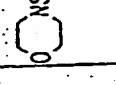


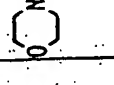
Compd. of Ex. No.	R ¹	R ²	R ³	X	R ⁴	[13] ([1A])	
						Yield%	NMR (δ)
28	m-(N-formyl)methyl- amino- phenyl			NH		91	0.60-1.78(13H, m), 2.51(2H, m), 2.84(4H, m), 3.17(2H, m), 3.37(3H, s), 3.40(5H, m), 3.54(1H, dd, J=4.1, 10Hz), 3.96(2H, m), 4.14(1H, m), 4.74(1H, m), 5.20(1H, d, J=5.7Hz), 6.60(1H, d, J=6.6Hz), 7.15(1H, d, J=1.9Hz), 7.34(5H, m), 7.53(1H, t, J=8.9Hz), 7.89(2H, m), 8.61(1H, s), 8.84(1H, d, J=2.1Hz), 9.26(1H, d, J=7.2Hz)
29	N-methyl- pyrrolyl			Cl ₂	+SO ₂	85	0.70-1.80(13H, m), 2.63(1H, dd, J=9.6, 16.8Hz), 2.80(1H, dd, J=2.5, 16.8Hz), 2.95(2H, m), 3.07-3.33(3H, m), 3.48(2H, m), 3.69(3H, s), 3.90(1H, ddd, J=7.4Hzx3), 4.06(1H, m), 4.75(1H, ddd, J=5.6Hz), 6.36(1H, d, J=9.6Hz), 6.57(1H, s), 6.58(1H, s), 7.11(1H, d, J=1.8Hz), 7.28(6H, m), 7.49(1H, d, J=6.8Hz), 8.69(1H, d, J=2Hz)
30	(N-morpho- lino)- methyl			NH		48	0.58-2.00(13H, m), 2.58(8H, m), 2.98(3H, m), 3.27(4H, m), 3.60(1H, dd, J=4.8, 14.8Hz), 3.80(6H, m), 4.15(2H, m), 4.78(1H, m), 5.03(1H, d, J=4.5Hz), 6.60(1H, d, J=9.4Hz), 7.18(1H, d, J=1.8Hz), 7.44(1H, s), 7.45(1H, ddd, J=7Hzx3), 7.61(1H, t, J=7.2Hz), 7.72(1H, t, J=7.0Hz), 7.86(1H, d, J=8.2Hz), 7.94(1H, d, J=7.0Hz), 8.42(1H, d, J=8.2Hz), 8.86(1H, d, J=1.8Hz), 9.45(1H, d, J=7.8Hz)
31	(N-pyperi- dino)- methyl			NH		58	0.60-2.08(19H, m), 2.54(7H, m), 2.71(2H, m), 2.99(3H, m), 3.22(1H, m), 3.28(2H, s), 3.60(1H, dd, J=5.15Hz), 3.80(2H, m), 4.14(2H, m), 4.80(1H, m), 5.07(1H, bs), 6.64(1H, d, J=8.6Hz), 7.18(1H, d, J=1.8Hz), 7.45(2H, ddd, J=7.0Hzx3), 7.60(1H, t, J=6.6Hz), 7.70(1H, t, J=6.6Hz), 7.86(1H, d, J=8.6Hz), 7.94(1H, d, J=7.8Hz), 8.24(1H, d, J=8.6Hz), 8.84(1H, d, J=1.8Hz), 9.34(1H, d, J=7.4Hz)
32	4-pyridyl			NH		95	0.78-1.68(13H, m), 2.45(2H, m), 2.75(3H, s), 2.83(3H, m), 3.10(3H, m), 3.43(4H, m), 3.45(2H, m), 3.97(2H, m), 4.12(1H, m), 4.75(1H, m), 5.16(1H, d, J=5Hz), 6.70(1H, d, J=9.4Hz), 6.92(1H, s), 7.33(5H, m), 7.82(2H, bs), 8.85(2H, bs), 9.22(1H, d, J=7.6Hz)

Table 9 (continued)

Compd. of Ex. No.	R ¹	R ²	R ³	X	R ⁴	Yield%	[13] ([1A])	
							NMR(δ)	
33	phenyl			Cl ₂	+SO ₂	71	0.70-1.85(13H, m), 1.30(9H, s), 2.84(1H, m), 3.14(7H, m), 3.46(1H, m), 4.06(1H, m), 4.13(1H, ddd, J=7Hz), 4.63(1H, m), 6.40(1H, d, J=10Hz), 6.79(1H, s), 7.25(5H, m), 7.48(2H, t, J=7.5Hz), 7.57(1H, m), 7.97(2H, m), 8.51(1H, m)	
34	4-pyridyl	-CONH ₂		NH		70	0.79-1.78(13H, m), 2.10(2H, m), 2.50(2H, m), 2.68(3H, m), 2.96(4H, m), 3.13(2H, m), 3.93(1H, dd, J=3.6, 14.2Hz), 4.05(1H, m), 4.21(2H, m), 4.78(1H, m), 7.28(1H, d, J=9.0Hz), 7.44(2H, d, J=4.4Hz), 7.58(2H, m), 7.88(4H, m), 8.20(1H, d, J=7.8Hz), 8.50(1H, d, J=8.0Hz), 8.80(2H, bs)	
35	4-pyridyl	-SMe		NH		84	0.80-1.80(13H, m), 2.14(3H, s), 2.63(2H, m), 2.96(1H, m), 3.26(3H, m), 3.49(4H, m), 3.98(1H, dddd, J=5.0Hz x 4), 4.12(1H, m), 4.25(1H, m), 4.65(1H, ddd, J=5.8Hz), 4.99(1H, d, J=5.4Hz), 6.90(1H, d, J=10Hz), 7.30(5H, m), 7.78(2H, bs), 8.83(2H, bs)	
36	4-pyridyl			Cl ₂	+SO ₂	89	0.78-1.80(13H, m), 2.95-3.55(10H, m), 3.99(1H, ddd, J=7.8Hz), 4.18(1H, m), 4.63(1H, ddd, J=5.6Hz), 6.39(1H, d, J=9.4Hz), 7.11(1H, d, J=1.8Hz), 7.38(2H, d, J=5.2Hz), 7.58(3H, m), 7.76(3H, m), 7.89(1H, m), 8.03(1H, d, J=7.8Hz), 8.54(1H, d, J=1.8Hz), 8.80(2H, bs)	
37	phenyl	TS 		Cl ₂		89	0.70-1.82(13H, m), 1.22(3H, d, J=7Hz), 1.30(3H, d, J=7Hz), 1.90(1H, bs), 2.43(3H, s), 2.72-3.23(8H, m), 3.49(1H, dd, J=13.2, 9.6Hz), 3.99(1H, m), 4.19(1H, m), 4.57(1H, ddd, J=6Hz x 3), 6.40(1H, d, J=9.4Hz), 7.96(2H, d, J=6.8Hz), 7.15-7.32(6H, m), 7.34(2H, d, J=8.4Hz), 7.42-7.63(3H, m), 7.85(2H, d, J=8.4Hz), 7.85(1H, d, J=8.4Hz)	
38	phenyl	TS 		Cl ₂		98	0.70-1.83(13H, m), 1.23(3H, t, J=7.4Hz), 2.43(3H, s), 2.20(2H, bs), 2.65-3.24(8H, m), 3.50(1H, dd, J=9.6, 14.2Hz), 4.00(1H, m), 4.19(1H, m), 4.57(1H, ddd, J=6.2Hz x 3), 6.43(1H, d, J=9.4Hz), 7.06-7.40(8H, m), 7.40-7.63(3H, m), 7.80(2H, d, J=8.4Hz), 7.88(1H, s), 7.97(2H, d, J=1.8Hz)	

Table 9 (continued)

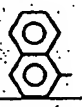


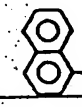
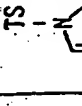

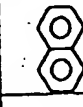

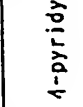
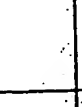
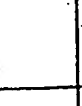
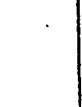
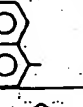

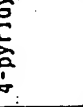
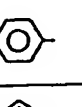
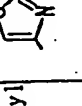

Compd. of Ex. No.	R ¹	R ²	R ³	X	R ⁴	[13] ([1A])	
						Yield%	NMR(δ)
39	phenyl			Cl ₂		76	0.70~1.80(13H, m), 2.28(1H, dd, J=6.2, 16.4Hz), 2.66(1H, dd, J=4.16, 8Hz), 2.92(3H, m), 3.05~3.75(12H, m), 4.06(1H, m), 4.19(1H, m), 4.79(1H, ddd, J=6.2Hz), 6.78(1H, d, J=10Hz), 7.18(1H, d, J=1.8Hz), 7.30~7.57(7H, m), 7.77(1H, d, J=7.8Hz), 7.88(1H, m), 8.04(3H, m), 8.29(1H, d, J=6.4Hz), 8.69(1H, d, J=1.8Hz), 0.80~1.80(13H, m), 2.40(3H, s), 2.90~3.75(17H, m), 4.06(1H, m), 4.22(1H, d, J=9.2Hz), 4.63(1H, ddd, J=5Hz), 6.87(1H, d, J=9.6Hz), 7.18~7.59(9H, m), 7.74~8.14(10H, m)
40	phenyl			Cl ₂		81	0.70~1.79(13H, m), 2.25(1H, dd, J=6.6, 17Hz), 2.74(1H, dd, J=3.8, 16.8Hz), 2.88(1H, dd, J=3.6, 16Hz), 2.96(3H, m), 3.08~3.64(10H, m), 3.75(1H, dd, J=4.8, 13Hz), 4.12(2H, m), 4.75(1H, ddd, J=5.8Hz), 6.82(1H, d, J=9.4Hz), 7.19(1H, d, J=2Hz), 8.77(2H, bs), 7.28(1H, d, J=7.6Hz), 7.41(1H, t, J=7Hz), 7.54(2H, m), 7.78(1H, d, J=8Hz), 7.87(3H, m), 8.03(1H, m), 8.46(1H, d, J=6.2Hz), 8.71(1H, d, J=2Hz)
41	4-pyridyl			Cl ₂		76	0.70~1.90(13H, m), 2.49(2H, m), 2.87(3H, m), 3.15(3H, m), 3.41(4H, m), 3.57(2H, m), 3.97(2H, m), 4.11(1H, m), 4.74(1H, m), 5.11(1H, d, J=5Hz), 6.59(1H, d, J=9.2Hz), 7.15(1H, d, J=2Hz), 7.32(5H, m), 7.85(2H, m), 8.85(1H, d, J=2Hz), 8.85(2H, m), 9.34(1H, d, J=7Hz)
42	4-pyridyl			Cl ₂		79	0.76~1.86(13H, m), 1.95(2H, m), 2.50(2H, m), 2.66(2H, m), 2.88~3.29(6H, m), 3.58(1H, dd, J=5.15Hz), 3.98(1H, m), 4.13(3H, m), 4.79(1H, m), 5.05(1H, d, J=4.5Hz), 6.63(1H, d, J=10Hz), 7.17(1H, d, J=1.7Hz), 7.43(2H, m), 7.59(2H, m), 7.88(4H, m), 8.69(1H, m), 8.85(1H, d, J=2Hz), 8.55(2H, bs), 9.51(1H, d, J=7Hz)
43	4-pyridyl			NH		95	
44	4-pyridyl			NH		96	

Table 9 (continued)



















Compd. of Ex. No.	R ¹	R ²	R ³	X	R ⁴	[13] ([1A])	
						Yield%	NMR(δ)
45	cyclohexyl			NH		94	0.77-1.85(23H, m), 2.36(1H, m), 2.56(4H, m), 2.85(3H, m), 3.18(1H, dd, J=5.2, 14.4 Hz), 3.49(6H, m), 3.90(3H, m), 4.75(1H, m), 5.01(1H, d, J=5.8 Hz), 6.55(1H, d, J=9.2 Hz), 7.16(1H, d, J=1.8 Hz), 7.32(5H, m), 8.84(1H, d, J=1.8 Hz), 9.10(1H, d, J=7.2 Hz)
46	4-pyridyl			NH		89	0.77-1.80(13H, m), 2.35(6H, s), 2.75(1H, dd, J=10.6, 13.8 Hz), 3.15(3H, m), 3.43(1H, dd, J=3.8, 13.8 Hz), 3.55(1H, dd, J=4.2, 15.1 Hz), 3.75(1H, m), 3.95(2H, m), 4.12(1H, m), 4.73(1H, m), 4.98(1H, d, J=5.1 Hz), 6.63(1H, d, J=9.4 Hz), 7.13(1H, d, J=2.1 Hz), 7.31(5H, m), 7.85(2H, bs), 8.84(1H, d, J=2.1 Hz), 8.84(2H, bs), 9.39(1H, d, J=7.2 Hz)
47	4-pyridyl			NH		90	0.79-1.80(13H, m), 2.30(2H, m), 2.42(2H, m), 2.85-3.62(11H, m), 3.94(1H, m), 4.00(1H, m), 4.20(1H, m), 4.42(1H, m), 4.75(1H, m), 6.63(1H, d, J=9.4 Hz), 7.16(1H, d, J=2.1 Hz), 7.29(5H, m), 7.71(1H, d, J=4.4 Hz), 7.85(2H, m), 8.33(1H, d, J=7.1 Hz), 8.62(1H, d, J=2.1 Hz), 8.83(2H, bs)
48	4-pyridyl			NH		94	0.79-1.76(13H, m), 2.20(2H, m), 2.49(2H, m), 2.80-3.08(4H, m), 3.19-3.56(7H, m), 3.92(1H, dd, J=4.4, 14.2 Hz), 4.00(1H, m), 4.20(1H, m), 4.55(1H, m), 4.74(1H, m), 6.65(1H, d, J=9.6 Hz), 7.15(1H, d, J=1.6 Hz), 7.45(2H, d, J=5.4 Hz), 7.57(2H, m), 7.85(4H, m), 8.19(1H, m), 8.33(1H, d, J=7.1 Hz), 8.52(1H, d, J=2.1 Hz), 8.83(2H, bs)
49	2-(N-morpholino)ethoxy-phenyl			NH		77	0.82-1.80(13H, m), 2.19(2H, m), 2.33(2H, m), 2.60(4H, m), 2.77-3.25(6H, m), 3.43(7H, m), 3.74(4H, m), 3.83(1H, dd, J=5.15 Hz), 3.98(1H, m), 4.21(3H, m), 4.63(1H, m), 4.76(1H, dddd, J=5.2 Hz x 4), 4.76(1H, m), 6.64(1H, d, J=9.8 Hz), 7.12(1H, d, J=1.4 Hz), 7.15(1H, m), 7.41(3H, m), 7.55(3H, m), 7.67(2H, m), 7.84(2H, m), 8.17(1H, m), 8.54(1H, d, J=2.1 Hz)
50	4-pyridyl			NH		83	0.60-1.80(13H, m), 3.08(5H, m), 3.33(4H, m), 3.57(2H, m), 3.64(4H, m), 4.05(1H, m), 4.18(1H, m), 4.65(1H, m), 4.95(1H, bs), 6.87(1H, d, J=9.8 Hz), 7.17(1H, d, J=1.2 Hz), 7.30(5H, m), 8.17(2H, m), 8.58(1H, d, J=9.1 Hz), 8.70(1H, d, J=1.2 Hz), 8.84(2H, m)

Table 9 (continued)


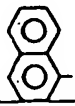
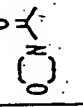

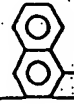
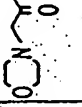
Compd. of Ex. No.	R ¹	R ²	R ³	X	R ⁴	[13] ([1A])	
						Yield%	NMR(δ)
51	m-2-(N-morpholino)ethoxy-phenyl			ClI ₂		70	0.70~1.80(13H, m), 2.28(1H, dd, J=6.6, 17.5Hz), 2.56(4H, m), 2.68(1H, dd, J=4.16, 6Hz), 2.80(3H, m), 2.95(1H, m), 3.18(2H, m), 3.35(4H, m), 3.50(2H, m), 3.63(1H, dd, J=5.4, 12Hz), 3.71(4H, t, J=4.4Hz), 3.87(1H, d, J=7.4Hz), 4.05(1H, m), 4.16(3H, t, J=5.6Hz), 4.77(1H, ddd, J=6.4Hzx3), 6.79(1H, d, J=9.6Hz), 7.09(1H, dd, J=2.4, 8Hz), 7.18(1H, d, J=1.8Hz), 7.35(3H, m), 7.54(3H, m), 7.63(1H, d, J=7.8Hz), 7.77(1H, d, J=7.6Hz), 7.88(1H, m), 8.02(1H, m), 8.28(1H, d, J=6.4Hz), 8.70(1H, d, J=1.8Hz)
52	cyclohexyl			Nil		94	0.78(23H, m), 2.22(2H, m), 2.38(2H, m), 2.58(2H, m), 3.00(3H, m), 3.48(6H, m), 3.82(1H, dd, J=5.14, 6Hz), 3.82(1H, m), 3.98(1H, m), 4.63(1H, m), 4.74(1H, ddd, J=8.8Hz), 6.56(1H, d, J=8.8Hz), 7.12(1H, d, J=1.8Hz), 7.44(2H, m), 7.58(2H, dt, J=1.8, 6.4Hz), 7.80(1H, m), 7.89(1H, m), 8.11(1H, d, J=7.8Hz), 8.22(1H, d, J=7.2Hz), 8.54(1H, d, J=1.8Hz)

Table 9 (continued)

Compd. of Ex. No.	$[\alpha]_D^{25}$ C=1.0, CHCl ₃ (Temp. °C)	Molecular formula (Molecular weight)	Calcd.	Found	IR ν_{\max} cm ⁻¹ CHCl ₃
23	-20.1 (24)	C ₃₇ H ₄₀ N ₃ O ₆ S ₂ · 1/2 H ₂ O (704.94)	C: 63.04 H: 7.15 N: 5.96 S: 9.10	C: 63.28 H: 7.21 N: 5.91 S: 8.97	3520, 3420, 3360 (br) 1670, 1600, 1580, 1450, 1118
24	-22.6 (24)	C ₃₆ H ₄₈ N ₄ O ₆ S ₂ · 1/4 H ₂ O (708.93)	C: 61.65 H: 6.77 N: 7.99 S: 9.14	C: 61.47 H: 7.02 N: 8.01 S: 8.91	3410, 3360, 1665, 1605, 1595, 1505, 1450, 1410, 1115
25	-23.1 (25)	C ₃₅ H ₄₇ N ₃ O ₆ S ₃ (701.948)	C: 59.89 H: 6.75 N: 5.99 S: 13.70	C: 59.68 H: 6.71 N: 5.89 S: 13.41	3315, 1665, 1510, 1412, 1290, 1115
26	-19.6 (24)	C ₃₇ H ₅₅ N ₃ O ₆ S ₂ · 1/2 H ₂ O (710.99)	C: 62.51 H: 7.94 N: 5.91 S: 9.02	C: 62.60 H: 8.05 N: 5.76 S: 8.87	3520, 3420, 3360, (br-sh), 1665, 1605, 1510, 1450, 1118
27	-14.4 (23.5)	C ₄₃ H ₆₀ N ₄ O ₆ S ₂ · 1/2 H ₂ O (834.101)	C: 61.92 H: 7.37 N: 6.72 S: 7.69	C: 61.77 H: 7.51 N: 6.52 S: 7.41	3500, 3420, 3360, 1665, 1596, 1581, 1505, 1460, 1448
28	(24.0) -23.2 (McOII)	C ₃₈ H ₅₀ N ₆ O ₆ S ₂ · 3/4 H ₂ O · 1/2 CH ₂ Cl ₂ (836.462)	C: 55.23 H: 6.30 N: 10.04 S: 7.66	C: 55.02 H: 6.07 N: 10.01 S: 7.44	3370, 1672, 1603, 1586, 1511, 1450, 1406, 1341, 1262, 1158, 1116
29	-24.1 (24.0)	C ₃₀ H ₅₀ N ₄ O ₆ S ₂ · 1/2 H ₂ O · 1/4 iPr ₂ O (733.490)	C: 61.41 H: 7.49 N: 7.64 S: 8.74	C: 61.28 H: 7.43 N: 7.50 S: 8.50	3500, 3420, 3360, 1660, 1605, 1530, 1508, 1462, 1450
30		C ₃₀ H ₅₄ N ₆ O ₆ S ₂ · 3/2 H ₂ O · 3/4 iPr ₂ O (902.655)	C: 57.88 H: 7.54 N: 9.31 S: 7.10	C: 57.65 H: 7.31 N: 9.59 S: 6.88	3380(3300), 1712, 1665, 1600(1535), 1510, 1455, 1430

Table 9 (continued)

Compd. of Ex. No.	$[\alpha]_D^{25}$ C=1.0, CHCl ₃ (Temp. °C)	Molecular formula (Molecular weight)	Calcd.	Found	IR ν max CHCl ₃ cm ⁻¹
31		C ₄₀ H ₅₀ N ₆ O ₇ S ₂ · 1/2 H ₂ O · 1/4 iPr ₂ O (840.587)	C: 59.30 H: 7.37 N: 10.00 S: 7.63	C: 59.44 H: 7.45 N: 9.88 S: 7.55	3380 (3280), 1705, 1662, 1600, 1535, 1510, 1450, 1400
32	-37.0 (24) 162-64	C ₃₀ H ₄₀ N ₆ O ₇ S ₂ · 1/4 H ₂ O (745.044)	C: 58.01 H: 6.56 N: 11.27 S: 8.60	C: 57.94 H: 6.53 N: 11.36 S: 8.36	3560, 3380 (3300), 1675, 1602, 1595, 1510 (1535), 1455
34		C ₃₇ H ₄₈ N ₆ O ₈ S · 1/2 H ₂ O (745.876)	C: 59.58 H: 6.62 N: 11.27 S: 4.30	C: 59.37 H: 6.66 N: 11.39 S: 4.25	3600, 3360, 1732, 1685, 1664, 1640, 1600, 1545
35	-62.6 (24) 175-8	C ₂₃ H ₄₇ N ₆ O ₇ S ₂ · 1/2 H ₂ O (698.886)	C: 56.71 H: 6.92 N: 10.02 S: 9.17	C: 56.60 H: 6.77 N: 9.87 S: 9.01	3600, 3380, 1670 (1690), 1600, (1515, 1525), 1500
36	-14.5 (24)	C ₄₃ H ₆₀ N ₄ O ₈ S ₂ · 3/2 H ₂ O · 1/2 iPr ₂ O (825.074)	C: 62.60 H: 7.33 N: 6.79 S: 7.77	C: 62.65 H: 7.13 N: 6.79 S: 7.59	3500, 3400, 3370, 1665, 1598, 1510, (sh, 1550, 1525)
39	98-100	C ₄₂ H ₅₀ N ₄ O ₈ S · 1/2 H ₂ O (749.952)	C: 67.45 H: 6.87 N: 7.49 S: 4.29	C: 67.47 H: 6.93 N: 7.54 S: 4.23	
41	-19.7 (25.5)	C ₄₁ H ₄₀ N ₅ O ₈ S · H ₂ O (757.947)	C: 64.97 H: 6.78 N: 9.24 S: 4.23	C: 65.08 H: 6.82 N: 9.21 S: 3.95	3400, 3340, 1665, (sh, 1695), 1625, 1530, 1510, 1450, 1410
43	-33.2 (25.5)	C ₃₅ H ₄₀ N ₆ O ₇ S ₂ (726.921)	C: 57.83 H: 6.38 N: 11.56 S: 8.82	C: 57.89 H: 6.36 N: 11.47 S: 8.72	3560, 3540, 3380, (sh, 3300), 1665, 1500, 1530, 1510

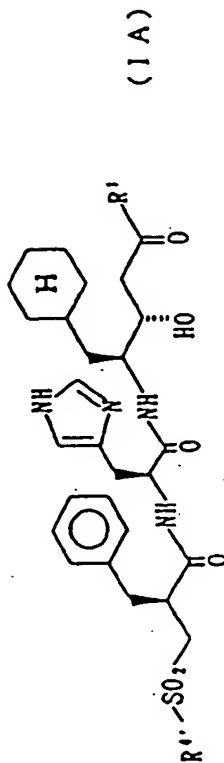
Table 9 (continued)

Compd. of	$[\alpha]_D^{25}$ C=1.0, CHCl ₃ (Temp. °C)	Molecular formula (Molecular weight)	Calcd.	Found	IR ν_{\max} CHCl ₃ cm ⁻¹
44	-40.7(24) 108-110	C ₄₁ H ₅₃ N ₂ O ₆ S ₂ (830.014)	C:59.33 H:6.44 N:10.13 S:7.73	C:59.40 H:6.50 N:10.04 S:7.50	3560, 3520, 3390, (sh 3300), 1670, 1600, 1535, 1510
45	-34.6 (24)	C ₃₀ H ₃₃ N ₅ O ₇ S · 3/4 H ₂ O · 1/2 iPr ₂ O (796.554)	C:58.80 H:7.78 N:8.79 S:8.05	C:58.55 H:7.55 N:9.05 S:7.77	3540, 3380, 3300, 1670, 1605, 1530, 1510, 1450, 1408
46	-32.0 (25)	C ₃₃ H ₄₄ N ₆ O ₈ S ₂ · 1/2 H ₂ O · 1/2 iPr ₂ O · 1/4 CHCl ₃ (766.200)	C:56.83 H:6.91 N:10.97 S:8.37	C:56.85 H:6.82 N:11.14 S:8.01	3550, 3380(3300), 1665, 1605, 1596, 1530, 1510, 1455, 1450
47		C ₃₇ H ₄₈ N ₆ O ₈ S · 3/2 H ₂ O · 1/10 iPr ₂ O (733.120)	C:60.71 H:7.03 N:11.46 S:4.37	C:60.79 H:7.05 N:11.56 S:4.31	
48	-23.0(25)	C ₄₁ H ₅₀ N ₆ O ₈ S · 5/4 H ₂ O (777.444)	C:68.34 H:6.81 N:10.81 S:4.12	C:63.34 H:6.92 N:10.64 S:3.81	3520, 3340, 1670, 1600, 1510(1530)
49	-19.3(25)	C ₄₈ H ₆₂ N ₆ O ₈ S · 3/2 H ₂ O (910.114)	C:63.34 H:7.20 N:9.23 S:3.52	C:63.49 H:7.27 N:9.35 S:3.36	3520, 3340, 1670, 1598, 1580, 1510, (sh 1530)
50	-24.8 (24)	C ₃₀ H ₄₀ N ₆ O ₈ S ₂ · H ₂ O · 1/5 D ₂ O (729.31)		C:61.35 H:7.03 N:11.36 S:4.16	3430(3480)3320, 1670, 1640, 1603, 1510

Table 9 (continued)

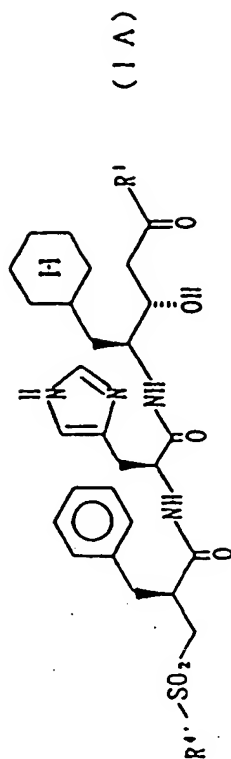
Compd. of	$[\alpha]_D^{25}$ C=1.0, MeOH (Temp. °C)	Molecular formula (Molecular weight)	Calcd.	Found	IR ν max cm ⁻¹
Ex. No. 51	-15.6 (25)	C ₄₈ H ₉₁ N ₅ O ₁₁ S (881.586)	C: 65.39 H: 7.15 N: 7.94 S: 3.64	C: 65.63 H: 7.44 N: 7.85 S: 3.39	3400, 3340, 1668, (1635), 1600, 1585, 1511, 1460, 1440
52	-26.0 (24)	C ₄₂ H ₅₇ N ₅ O ₁₁ S (768.990)	C: 65.60 H: 7.60 N: 9.11 S: 4.17	C: 65.66 H: 7.68 N: 9.08 S: 3.89	3480, 3340, 1670, 1598, 1508, (1525), 1448, 1425, 1410

Table 10



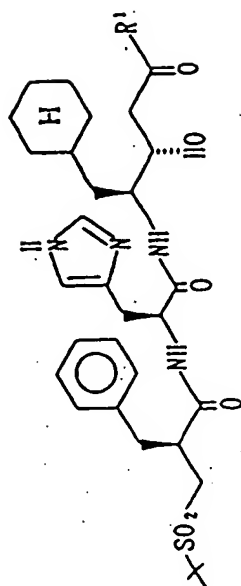
Compd. of Ex. No.	R ¹	R ²	Yield %	[α] _D ²⁰ (C=1.0, MeOH) (°C)	Molecular formula	Elemental analysis		IR ν _{max} cm ⁻¹
						Calcd.	Found	
2	phenyl	tert-butyl	75	-22.2° (24°C)	C ₂₁ H ₂₅ N ₂ O ₄ S ·1/2H ₂ O, 1/4iPr ₂ O	C: 64.82 H: 7.70 N: 7.85 S: 4.49	C: 64.87 H: 7.65 N: 7.99 S: 4.33	3460, 3360(br), 1663, 1600, 1580, 1498, 1450, 1116
3	o-fluorophenyl	tert-butyl	90	-20.9 (24.0)	C ₂₁ H ₁₉ FN ₂ O ₄ S ·3/4H ₂ O	C: 62.56 H: 7.17 N: 7.89 S: 4.51	C: 62.65 H: 7.12 N: 8.05 S: 4.56	3460, 3360(br), 1666, 1611, 1575, 1480, 1453, 1116
4	m-methoxyphenyl	tert-butyl	73	-18.1 (24.5)	C ₂₁ H ₂₃ N ₂ O ₄ S ·1.5H ₂ O	C: 62.02 H: 7.53 N: 7.61 S: 4.36	C: 62.05 H: 7.16 N: 7.52 S: 4.12	3468, 3348(br), 1668, 1600, 1585, 1499, 1464, 1451, 1430, 1289, 1258, 1169, 1117, 1077.
5	p-methylphenyl	tert-butyl	94	-21.3 (24.0)	C ₂₁ H ₂₃ N ₂ O ₄ S ·3/4H ₂ O	C: 64.61 H: 7.63 N: 7.93 S: 4.54	C: 64.65 H: 7.64 N: 7.99 S: 4.61	3460, 3350(br), 1666, 1608, 1564, 1498, 1450, 1116
6	2,4-difluorophenyl	tert-butyl	93	-21.0 (23.5)	C ₂₁ H ₁₉ F ₂ N ₂ O ₄ S ·1/2H ₂ O	C: 61.39 H: 6.82 N: 7.74 S: 4.43	C: 61.18 H: 6.82 N: 7.76 S: 4.40	3470, 3350(br), 1665, 1612(1595sh), 1498, 1450, 1430, 1116, 1100, 970, 855

Table 10 (continued)



Compd. of Ex. No.	R'	R''	Yield (C=1.0, MeOH) %	[α] _D ²⁰ (C=1.0, MeOH) (°C)	Molecular formula	Elemental analysis		IR ν _{max} cm ⁻¹
						Calcd.	Found	
7	1-naphthyl	tert-butyl	60	-15.3 (24.0)	C ₂₁ H ₁₅ N ₃ O ₆ S ·5/4H ₂ O	C: 65.53 H: 7.31 N: 7.46 S: 4.27	C: 65.43 H: 7.09 N: 7.37 S: 4.24	3672, 3352(br), 1665, 1605, 1509, 1464, 1450, 1441, 1369, 1291, 1117, 1077
8	3-thienyl	tert-butyl	94	-24.4 (28.0)	C ₂₃ H ₁₆ N ₃ O ₆ S ₂ ·H ₂ O	C: 59.81 H: 7.17 N: 7.97 S: 9.12	C: 59.68 H: 7.02 N: 8.23 S: 9.10	3460, 3350(br), 1662, 1600, 1505, 1446, 1113
9	2-thiazolyl	tert-butyl	71		C ₂₄ H ₁₇ N ₃ O ₆ S ₂ ·2H ₂ O · 1/2C ₄ H ₈ O ₃	C: 56.45 H: 7.24 N: 9.16 S: 8.37	C: 56.41 H: 6.95 N: 9.14 S: 8.09	3660, 3356(br), 1661, 1599, 1511, 1446, 1113

Table 10 (continued)

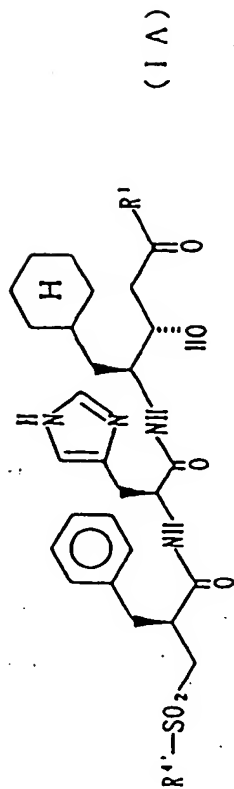


Compd. of Ex. No.	R ¹	Yield %	[α] _D ²⁰ C=1.0, MeOH (°C)	Molecular formula	Calcd.	Found	I R ν _{max} cm ⁻¹
10	m-fluorophenyl	88	-21.4 (24.0)	C ₂₇ H ₁₉ N ₃ FO ₃ S 3/4H ₂ O	C: 62.56 F: 2.67 H: 7.17 N: 7.89 S: 4.51	C: 62.72 F: 2.60 H: 7.14 N: 7.67 S: 4.60	3470, 3340, 1663, 1605, 1590, 1496, 1445, 1400, 1370, 1115, 1075, 1015
11	p-fluorophenyl	87	-21.3 (24.0)	C ₂₇ H ₁₉ N ₃ FO ₃ S 3/4H ₂ O	C: 62.56 F: 2.67 H: 7.17 N: 7.89 S: 4.51	C: 62.60 F: 2.66 H: 7.17 N: 7.84 S: 4.62	3470, 3340, 1665, 1600, 1505, 1450, 1410, 1370, 1156, 1115, 1076
12	2,6-difluoro- phenyl	75	-23.4 (23.5)	C ₂₇ H ₁₉ N ₃ F ₂ O ₃ S 2/3H ₂ O	C: 61.15 F: 5.23 H: 6.84 N: 7.71 S: 4.41	C: 61.05 F: 5.26 H: 6.60 N: 7.77 S: 4.75	3468, 3360, 1664, 1624, 1502, 1467, 1450, 1402, 1370, 1290, 1117, 1078, 1016, 991

Table 10 (continued)

Compd. of Ex. No.	R'	Yield %	$[\alpha]_D^{25}$ C=1.0, MeOH (°C)	Molecular formula	Calcd.	Found	I R ν_{\max} cm^{-1}
13	<i>o</i> -methoxyphenyl	75	-3.3 (24.0)	$\text{C}_{10}\text{H}_{11}\text{N}_2\text{O}_4\text{S}$ $\cdot 3/4\text{H}_2\text{O}$	C: 63.18 H: 7.46 N: 7.76 S: 4.44	C: 63.18 H: 7.52 N: 7.38 S: 4.03	3468, 3348, 1665, 1600, 1502, 1487, 1465, 1438, 1289, 1163, 1117, 1077, 1026
14	<i>o</i> -chlorophenyl	75	-8.4 (23.5)	$\text{C}_{10}\text{H}_9\text{ClN}_2\text{O}_4\text{S}$ $\cdot 0.1 \text{ Cl}_2\text{Ce}_2 \cdot 1/3\text{H}_2\text{O}$	C: 61.23 Cl: 5.85 H: 6.91 N: 7.70 S: 4.41	C: 61.03 Cl: 5.63 H: 6.85 N: 7.68 S: 4.31	3468, 3360, 1665, 1593, 1500, 1450, 1434, 1402, 1370, 1291, 1117, 1077
15	<i>o</i> -cyanophenyl	92	-20.6 (24.5)	$\text{C}_{10}\text{H}_8\text{N}_4\text{O}_4\text{S}$ $\cdot 1/2\text{O} \cdot 1/4\text{Cl}_2\text{Ce}_2$	C: 61.82 H: 6.99 N: 9.42 S: 4.31	C: 61.87 H: 6.75 N: 9.40 S: 4.25	3468, 3360, 2236, 1666, 1602, 1499, 1450, 1431, 1401, 1370, 1288, 1150, 1117, 1077, 909
16	<i>o</i> -methylsulfonyl- aminophenyl	86	-11.4 (24.0)	$\text{C}_{10}\text{H}_{11}\text{N}_3\text{O}_4\text{S}_2$ $\cdot 1/4(\text{iPr})_2\text{O} \cdot 1/2\text{H}_2\text{O}$	C: 58.82 H: 7.19 N: 8.68 S: 7.95	C: 58.54 H: 7.06 N: 8.46 S: 7.71	3464, 3352, 1664, 1607, 1578, 1492, 1452, 1400, 1340, 1289, 1155, 1117, 1077, 968, 917
17	<i>p</i> -trifluoroacetyl- phenyl	95	-18.8 (24.0)	$\text{C}_{10}\text{H}_9\text{F}_3\text{N}_2\text{O}_4\text{S}$ $\cdot 1/2\text{H}_2\text{O}$	C: 60.38 F: 7.54 H: 6.67 N: 7.41	C: 60.27 F: 7.53 H: 6.77 N: 7.27 S: 4.24	3450, 3350, 1665, 1605, 1510, 1450, 1410, 1325, 1170, 1135, 1115, 1065 S: 4.41

Table 10 (continued)



Compd. of Ex. No.	R'	R''	Yield %	[α] _D ²⁰ (C=1.0, MeOH)	Molecular formula	Calcd.	Found	IR ν _{max} cm ⁻¹
18	m-morpholino-carbonyloxyphenyl	tert-butyl	92	-15.8 ± 0.6 (25°C)	C ₂₃ H ₂₇ N ₃ O ₆ S · 1 3/4 H ₂ O	C: 60.09 H: 7.26 N: 8.34 S: 3.82	C: 59.93 H: 6.94 N: 8.38 S: 3.79	3470, 3320, 1711, 1680, 1665, 1605, 1587, 1500, 1420, 1370, 1116, 1068
19	m-morpholino-carbonylphenyl	tert-butyl	79	-15.8 ± 0.6 (25°C)	C ₂₃ H ₂₇ N ₃ O ₆ S · 5/4 H ₂ O	C: 61.93 H: 7.36 N: 8.60 S: 3.94	C: 61.70 H: 7.10 N: 8.42 S: 3.92	3356, 1665, 1627, 1581, 1498, 1463, 1451, 1369, 1289, 1117, 1075, 1028
20	3,4-methylenedioxypheyl	tert-butyl	88	-16.8 ± 0.6 (25°C)	C ₂₃ H ₂₇ N ₃ O ₆ S · 2H ₂ O · 2/5 dioxane	C: 59.89 H: 7.26 N: 7.05 S: 4.04	C: 59.86 H: 6.92 N: 6.77 S: 3.87	3470, 3340, 1665, 1605, 1505, 1490, 1445, 1117, 1080, 1042

Table 10 (continued)

Compd. of Ex. No.	R'	R''	Yield [α] (C-1.0. MeOH)	Molecular formula	Calcd.	Found	IR ν_{\max} cm ⁻¹
21	cyclohexyl	tert-butyl	94 -21.2 \pm 0.6 (24 \pm)	C ₃₇ H ₅₆ N ₄ O ₆ S -1/2H ₂ O	C: 62.42 H: 8.35 N: 7.87 S: 4.50	C: 62.42 H: 8.06 N: 8.10 S: 4.33	3460, 3340, 1662 (sh), 1685, 1605, 1500, 1450, 1115
22	p-methoxyphenyl	tert-butyl	91 -16.9 \pm 0.6 (24 \pm)	C ₃₈ H ₅₂ N ₄ O ₇ S -2.5H ₂ O - 1/2 dioxan	C: 60.21 H: 7.71 N: 7.02 S: 4.02	C: 60.24 H: 7.38 N: 6.78 S: 3.77	3470, 3340, 1665, 1603, 1575, 1510, 1170, 1116, 1076, 1030
37	phenyl	isopropyl	93 -21.2 \pm 0.6 (24 \pm)	C ₃₈ H ₄₈ N ₄ O ₆ S -H ₂ O	C: 63.32 H: 7.38 N: 8.20 S: 4.69	C: 63.59 H: 7.46 N: 8.01 S: 4.48	3460, 3340, 1665, 1600, 1580, 1500, 1450, 1120.
38	phenyl	ethyl	89 -22.1 \pm 0.6 (24 \pm)	C ₃₈ H ₄₈ N ₄ O ₆ S -3/4H ₂ O	C: 63.28 H: 7.21 N: 8.43 S: 4.83	C: 63.17 H: 7.18 N: 8.49 S: 4.60	3400, 3350, 1665, 1600, 1582, 1510, 1450, 1310, 1123

[illegible]78

Table 10 (continued)

Compd. of Ex. No.	NMR (δ):
2	0.70~1.83(13H, m), 1.33(9H, s), 2.68~3.18(7H, m), 3.27(1H, m), 3.60(1H, dd, J=9.8, 13.2Hz), 3.80(1H, br), 3.99(1H, m), 4.21(1H, m), 4.60(1H, m), 6.48(1H, d, J=9.4Hz), 6.89(1H, s), 7.08~7.63(8H, m), 7.50(1H, d, J=1.8Hz), 7.93(2H, d, J=8.4Hz)
3	0.68~1.83(13H, m), 1.33(9H, s), 2.70~3.17(7H, m), 3.60(1H, dd, J=13.1, 9.9Hz), 3.59(2H, brs), 3.25(1H, m), 3.98(1H, m), 4.18(1H, m), 4.59(1H, ddd, J=6.8, 6.8, 6.8Hz), 6.51(1H, d, J=9.0Hz), 6.90(1H, s), 7.05~7.37(7H, m), 7.52(1H, m), 7.55(1H, s), 7.84(1H, ddd, J=7.7, 1.9Hz)
4	0.70~1.83(13H, m), 1.32(9H, s), 2.74~3.17(7H, m), 3.27(1H, m), 3.59(1H, dd, J=10.13Hz), 3.83(3H, s), 4.00(1H, m), 4.20(1H, m), 4.60(1H, ddd, J=7.7, 7.7Hz), 4.72(1H, bs), 6.60(1H, d, J=9Hz), 6.86(1H, s), 7.04~7.55(9H, m), 7.48(1H, s)

Table 10 (continued)

Compd. of Ex. No.	NMR (δ):
5	0.70~1.85(13H, m), 1.32(9H, s), 2.40(3H, s), 2.70~3.18(7H, m), 3.27(1H, m), 3.61(1H, dd, J=13, 2, 9, 8Hz), 3.75(2H, bs), 3.99(1H, m), 4.19(1H, m), 4.61(1H, ddd, J=6, 5, 6, 5, 6, 5Hz), 6.53(1H, d, J=9, 11Hz), 6.88(1H, s), 7.10~7.45(7H, m), 7.52(1H, s), 7.82(2H, d, J=8, 2Hz)
6	0.70~1.80(13H, m), 1.34(9H, s), 2.77~3.16(7H, m), 3.20(2H, bs), 3.25(1H, m), 3.59(1H, dd, J=13, 9.5Hz), 3.48(1H, m), 4.18(1H, m), 4.59(1H, ddd, J=7, 7, 7Hz), 6.46(1H, d, J=9, 3Hz), 6.78~7.00(2H, m), 6.91(1H, s), 7.23(5H, m), 7.54(1H, s), 7.91(1H, ddd, J=8, 6, 8, 6, 6Hz)
7	0.70~1.85(13H, m), 1.30(9H, s), 2.82(1H, dd, J=13, 2, 8, 4Hz), 2.90~3.18(6H, m), 3.28(1H, m), 3.59(1H, dd, J=13, 0, 10, 0Hz), 4.03(1H, m), 4.24(1H, m), 4.60(1H, ddd, J=5, 1Hz), 5.30(1H, brs), 6.63(1H, d, J=9, 2Hz), 6.83(1H, s), 7.20(5H, m), 7.50(4H, m), 7.90(2H, m), 7.86(1H, s), 8.60(1H, d, J=7, 6Hz)
8	0.70~1.85(13H, m), 1.33(9H, s), 2.77~3.18(7H, m), 3.26(1H, m), 3.59(1H, dd, J=13, 2, 10, 2Hz), 3.97(1H, m), 4.18(1H, m), 4.58(1H, m), 6.44(1H, d, J=9, 4Hz), 6.88(1H, s), 7.12~7.47(6H, m), 7.50(1H, s), 7.51(1H, dd, J=5, 0, 1, 2Hz), 8.13(1H, dd, J=1, 2, 3Hz)
9	0.70~1.83(13H, m), 1.34(9H, s), 2.74~3.32(8H, m), 3.61(1H, dd, J=12, 8, 9, 4Hz), 3.73(2H, bs), 3.95(1H, m), 4.23(1H, m), 4.60(1H, ddd, J=6, 6, 6, 6Hz), 6.51(1H, d, J=9, 2Hz), 6.96(1H, s), 7.12~7.35(5H, m), 7.57(1H, s), 7.68(1H, d, J=3Hz), 8.00(1H, d, J=3Hz)

Table 10 (continued)

Compd. of Ex. No.	(1A)	
	NMR (δ)	
10	0.70~1.82(13H, m), 1.33(9H, s), 2.75~3.19(7H, m), 3.28(1H, m), 3.62(1H, dd, J=13, 10Hz), 4.02(1H, m), 4.20(1H, m), 4.59(1H, ddd, J=6, 4, 6, 4Hz), 6.55(1H, d, J=9, 4Hz), 6.92(1H, s), 7.25(6H, m), 7.45(2H, m), 7.56(1H, s), 7.62(1H, m), 7.70(1H, d, J=7, 6Hz)	
11	0.70~1.82(13H, m), 1.33(9H, s), 2.75~3.20(7H, m), 3.28(1H, m), 3.57(2H, bs), 3.60(1H, dd, J=13, 10Hz), 4.00(1H, m), 4.19(1H, m), 4.59(1H, ddd, J=6, 6, 6, 6Hz), 6.54(1H, d, J=9, 2Hz), 6.90(1H, s), 7.14(2H, dd, J=17, 2, 8, 8Hz), 7.26(6H, m), 7.48(1H, d, J=8Hz), 7.54(1H, s), 7.96(2H, dd, J=9, 5, 4Hz)	
12	0.70~1.85(13H, m), 1.33(9H, s), 2.78~3.18(7H, m), 3.25(1H, m), 3.61(1H, dd, J=13, 2, 9, 8Hz), 3.95(1H, m), 4.12(1H, m), 4.58(1H, ddd, J=6, 4, 6, 4Hz), 5.68(2H, bs), 6.55(1H, d, J=9, 4Hz), 6.85(1H, s), 6.93(2H, t, J=8, 2Hz), 7.15~7.48(7H, m), 7.49(1H, s)	
13	0.70~1.82(13H, m), 1.33(9H, s), 2.75~3.35(8H, m), 3.62(1H, dd, J=14, 10Hz), 3.87(3H, s), 3.95(1H, m), 4.15(1H, m), 4.62(1H, ddd, J=6, 5, 6, 5, 6, 5Hz), 6.66(1H, d, J=9, 4Hz), 6.82(1H, s), 6.85(2H, m), 7.24(5H, m), 7.46(1H, s), 7.46(1H, td, J=7, 8, 1, 8Hz), 7.70(1H, dd, J=7, 8, 1, 8Hz)	

Table 10 (continued)

Compd. of Ex. No.	(1A)	
	NMR (δ)	
14	0.70~1.80(13H, m), 1.33(9H, s), 2.78~3.17(7H, m), 3.26(1H, m), 3.52(1H, dd, J=13.9, 8Hz), 3.97(1H, m), 4.13(1H, m), 4.57(1H, ddd, J=6.7, 6.7, 6.7Hz), 6.52(1H, d, J=9.0Hz), 6.85(1H, s), 7.14~7.42(8H, m), 7.46(1H, s), 7.53(1H, m).	
15	0.70~1.80(13H, m), 1.33(9H, s), 2.77~3.18(7H, m), 3.29(1H, m), 3.61(1H, dd, J=12.8, 9.8Hz), 4.04(1H, m), 4.22(1H, m), 4.57(1H, ddd, J=5.8, 5.8, 5.8Hz), 6.59(1H, d, J=9Hz), 6.91(1H, s), 7.23(5H, m), 7.56(1H, s), 7.58(1H, t, J=7.8Hz), 7.81(1H, d, J=7.8Hz), 8.14(1H, d, J=8Hz), 8.23(1H, s).	
16	0.70~1.82(13H, m), 1.33(9H, s), 2.75~3.17(7H, m), 3.26(1H, m), 3.59(1H, dd, J=13.10, 4Hz), 4.01(1H, m), 4.16(1H, m), 4.57(1H, ddd, J=6.4, 6.4, 6.4Hz), 6.59(1H, d, J=9.2Hz), 6.88(1H, s), 7.20(6H, m), 7.49(1H, d, J=1.2Hz), 7.50(2H, m), 7.69(1H, dd, J=8.4, 1.2Hz), 7.86(1H, dd, J=8.2, 1.2Hz).	
17	0.70~1.80(13H, m), 1.32(9H, s), 2.75~3.18(7H, m), 3.27(1H, m), 3.58(1H, dd, J=13.4, 10Hz), 4.04(1H, m), 4.21(1H, t, d, J=7.2, 5Hz), 4.57(1H, ddd, J=6.3, 6.3, 6.3Hz), 6.54(1H, d, J=9.4Hz), 7.25(5H, m), 7.52(1H, d, J=9Hz), 7.71(2H, d, J=8.2Hz), 8.04(2H, d, J=8Hz).	





Table 10 (continued)

Compd of Ex. No.	(I A)	
	NMR (δ)	
18	0.70~1.83(13H, m), 1.34(9H, s), 2.70~3.15(7H, m), 3.28(1H, m), 3.48~3.82(8H, m), 3.88(1H, m), 4.14(1H, m), 4.63(1H, ddd, J=6.6, 6.6, 6.6 Hz), 6.39(1H, d, J=8.6 Hz), 6.80(1H, s), 7.17~7.40(6H, m), 7.40~7.60(3H, m), 7.77(1H, d, J=7.6 Hz)	
	0.70~1.80(13H, m), 1.33(9H, s), 2.63~3.17(7H, m), 3.30(1H, m), 3.38~3.95(10H, m), 4.18(1H, m), 4.63(1H, ddd, J=6.6, 6.6 Hz), 6.49(1H, d, J=8.8 Hz), 6.86(1H, s), 7.23(6H, m), 7.56(1H, s), 7.55(1H, m), 7.71(1H, d, J=6.6 Hz), 7.84(1H, s), 8.00(1H, m)	
	0.70~1.82(13H, m), 1.34(9H, s), 2.78~3.18(7H, m), 3.26(1H, m), 3.60(1H, ddd, J=13.9, 8.1 Hz), 3.97(1H, m), 4.17(1H, m), 4.60(1H, ddd, J=6.4, 6.4, 6.4 Hz), 6.04(2H, s), 6.45(1H, d, J=9.4 Hz), 6.84(2H, d, J=8.2 Hz), 6.86(1H, s), 7.25(5H, m), 7.39(1H, d, J=1.2 Hz), 7.49(1H, s), 7.51(2H, d, J=8.2 Hz)	

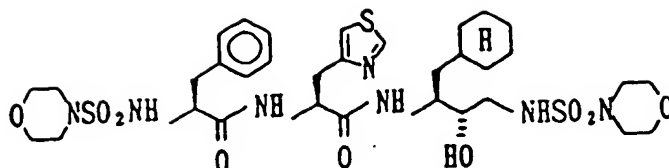
Table 10 (continued)

Compd. of Ex. No.	(I A)	
	NMR (δ)	
21	0.70~1.92(23H, m), 1.35(9H, s), 2.36(1H, m), 2.55(2H, m), 2.75~3.18(5H, m), 3.26(1H, m), 3.60(1H, dd, J=13, 3, 9, 8Hz), 3.88(1H, m), 3.99(1H, m), 4.56(1H, ddd, J=6, 6, 6Hz), 6.41(1H, d, J=9, 2Hz), 6.88(1H, s), 7.27(5H, m), 7.54(1H, s)	
	0.70~0.83(13H, m), 1.34(9H, s), 2.75~3.17(7H, m), 3.15(1H, m), 3.60(1H, dd, J=13, 9, 6Hz), 3.87(3H, s), 3.98(1H, m), 4.18(1H, m), 4.61(1H, ddd, J=6, 8, 6, 8Hz), 6.44(1H, d, J=9, 2Hz), 6.86(1H, s), 6.93(2H, d, J=9Hz), 7.25(5H, m), 7.49(1H, s), 7.91(2H, d, J=8, 6Hz)	
	0.70~1.85(13H, m), 1.23(3H, d, J=6, 8Hz), 1.30(3H, d, J=6, 8Hz), 2.70~3.18(8H, m), 3.22(1H, m), 3.56(1H, dd, J=13, 4, 9, 6Hz), 4.01(1H, m), 4.20(1H, m), 4.60(1H, ddd, J=6, 6Hz), 6.49(1H, d, J=9, 2Hz), 6.87(1H, s), 7.22(5H, m), 7.50(4H, m), 7.93(2H, d, J=6, 8Hz)	
38	0.70~1.85(13H, m), 1.23(3H, t, J=7, 4Hz), 2.65(8H, m), 3.23(1H, m), 3.57(1H, dd, J=14, 9, 8Hz), 4.03(1H, m), 4.19(1H, m), 4.62(1H, ddd, J=6, 2, 6, 2Hz), 6.56(1H, d, J=9, 2Hz), 6.88(1H, s), 7.20(5H, m), 7.45(1H, d, J=1, 6Hz), 7.52(3H, m), 7.93(2H, d, J=6, 8Hz)	

Table 10 (continued)

Compd. of Ex. No.	R ³	X	[I A]	
			NMR (δ)	
28		NH	0.65~1.77(13H, m), 2.50(2H, m), 2.80(3H, s), 3.00(2H, m), 3.20(1H, dd, J=5.15Hz), 3.40(4H, m), 3.53(1H, dd, J=5.15Hz), 3.95(2H, m), 4.12(1H, m), 4.80(1H, dt, J=4.7, 0.1Hz), 5.18(1H, d, J=6.1Hz), 6.62(1H, d, J=9.4Hz), 6.82(1H, dt, J=2.2, 7.1Hz), 7.14(1H, d, J=2.1Hz), 7.30(8H, m), 8.81(1H, d, J=2.1Hz), 9.09(1H, d, J=6.8Hz)	
33		Cl ₂	0.70~1.82(13H, m), 1.31(9H, s), 2.83~3.30(9H, m), 3.47(1H, dd, J=4.6, 13.2Hz), 4.01(1H, m), 4.16(1H, dt, J=3.6, 2.1Hz), 4.60(1H, ddd, J=4.8Hzx3), 5.30(1H, bs), 6.24(1H, s), 6.49(1H, d, J=9.8Hz), 7.25(5H, m), 7.47(2H, t, J=7.4Hz), 7.56(1H, d, J=7.1Hz), 7.62(1H, d, J=6.8Hz), 7.96(2H, dd, J=1.4, 6.6Hz)	
40		Cl ₂	0.78~1.80(13H, m), 2.56(2H, m), 2.90~3.70(15H, m), 4.04(1H, m), 4.21(1H, m), 4.64(1H, ddd, J=6.2Hz), 6.68(1H, d, J=10.1Hz), 6.87(1H, s), 7.26~7.59(8H, m), 7.76(1H, d, J=8.2Hz), 7.87(1H, m), 8.02(3H, m)	
42		Cl ₂	0.67~1.77(13H, m), 2.47(1H, dd, J=7.5, 17.1Hz), 2.62(1H, dd, J=5.18Hz), 2.88~3.76(15H, m), 4.08(1H, m), 4.17(1H, m), 4.63(1H, ddd, J=5.1Hz), 6.77(1H, d, J=10.1Hz), 6.88(1H, s), 7.29(1H, d, J=7.5Hz), 7.40(1H, t, J=7.5Hz), 7.54(3H, m), 7.80(2H, d, J=6.2Hz), 7.80(2H, m), 8.04(1H, m), 8.33(1H, m), 8.76(2H, d, J=6.1Hz)	

Examples 53



To the compound [25a] (24.5g, 41.6mmol) are added anisole (89.7g, 20eq) and anhydrous dichloromethane (250ml). To the mixture is dropwise added trifluoroacetic acid (250ml) with stirring and ice-cooling over 30 minutes, and the mixture is stirred at room temperature for one hour. The reaction mixture is concentrated *in vacuo*, made alkaline with Na_2CO_3 and saturated aqueous sodium bicarbonate, and extracted with a mixture of dichloromethane and methanol (9:1). The organic layer is washed with water, dried over MgSO_4 , and evaporated to dryness *in vacuo*. The residue is subjected to silica gel chromatography (SiO_2 : 600g, CH_2Cl_2 :MeOH: NH_4OH = 90:10:1) to obtain the compound [26a] (14.63g, 72%).

To the above compound [26a] (11.04g, 22.5mmol) are added N-(morpholin-4-ylsulfonyl)phenylalanine [12a] (8.5g, 1.2eq), HOBt (3.96g, 1.25eq), and anhydrous CH_3CN (200ml). To the mixture is added DCC (6.05g, 1.3eq) with stirring and ice-cooling, and the mixture is stirred at 0°C for one hour and then at room temperature for an additional one hour. To the reaction mixture is added ethyl acetate and it is then filtered. The filtrate is concentrated *in vacuo* and subjected to silica gel chromatography (SiO_2 : 600g, CH_2Cl_2 :MeOH = 97:3). Relevant fractions are combined and treated with isopropyl ether to give the compound [1b] (16.33g, 92%).

Elemental analysis (as $\text{C}_{33}\text{H}_{51}\text{N}_7\text{O}_9\text{S}_3 \cdot 0.75\text{H}_2\text{O} \cdot 1.0\text{CH}_2\text{Cl}_2$)

Calcd.: C: 49.20; H: 6.57; N: 12.13; S 11.90

Found : C: 49.05; H: 6.20; N: 11.92; S 11.78

$[\alpha]_D^{25} = -22.5$ ($c=1$; MeOH; 24°C)

IR: 3370, 2720, 1665, 1530, 1510, 1454, 1340, 1330, 1260, 1155, 1113, 1073, 943

NMR(δ): 0.72(3H,m), 1.12(6H,m), 4.16(1H,bd,J=8Hz), 1.62(3H,bd,J=8Hz), 2.21(1H,bs), 2.47(2H,m), 2.74 (1H, dd,J=10.14Hz), 2.80-3.33(4H,m), 3.21(4H,m), 3.33-3.62(8H,m), 3.75(4H,m), 3.97(2H,m), 4.68(1H,m), 5.16 (1H, d,J=5.4Hz), 5.64(1H,t,J=6.8Hz), 6.55(1H,d,J=9.2Hz), 7.19(1H,d,J=1.2Hz), 7.35(5H,m), 8.90(1H,d,J=1.2Hz), 9.40(1H,d,J=6.8Hz)

Examples 54-71

In accordance with substantially the same procedure as disclosed in Example 53, the compounds of the invention listed in Table 11 are obtained.

Table 11 (continued)

Compd. of Ex. No.	R ¹	R ²	R ³	X	R ⁴	Yield%	[α] _D ²⁰ C=1, MeOH(°C)	Molecular formula	[IB] Elemental analysis			IR ν _{max} cm ⁻¹
									Calcd.	Found		
59				NH	+ SO ₂ -	86	-17.6 (25.0)	C ₁₆ H ₁₃ N ₃ O ₂ S ₂ ·0.5H ₂ O ·0.25Cl ₂ ·Cf ₂	C: 57.81 H: 6.33 N: 11.72 S: 7.67	C: 57.84 H: 6.32 N: 11.69 S: 7.55	3330, 2920, 1670, 1599, 1575, 1580, 1505, 1336, 1165, 1115	
60				Cl ₂	+ SO ₂ -	70	-15.0 (23.5)	C ₁₈ H ₁₃ N ₃ O ₂ S ₂ ·0.2H ₂ O	C: 58.43 H: 6.46 N: 8.74 S: 12.00	C: 58.20 H: 6.27 N: 8.59 S: 11.75	3360, 2920, 1615, 1600, 1567, 1530, 1500, 1450, 1330, 1290	
61				Cl ₂	+ SO ₂ -	83	-3.4 (23.5)	C ₁₈ H ₁₃ N ₃ O ₂ S ₄ ·0.75H ₂ O	C: 53.28 H: 6.51 N: 7.31 S: 16.73	C: 53.04 H: 6.22 N: 7.56 S: 16.63	3360, 2920, 1660, 1530, 1510, 1450, 1405, 1338, 1290, 1155, 1115, 1015	
62				Cl ₂	+ SO ₂ -	72	-3.8 (25.5)	C ₁₈ H ₁₃ N ₃ O ₂ S ₃ ·0.5H ₂ O	C: 57.19 H: 6.80 N: 7.41 S: 12.72	C: 57.01 H: 6.69 N: 7.41 S: 12.81	3360, 2920, 1660, 1605, 1530, 1510, 1446, 1330, 1290, 1160, 1165, 1145, 1116, 1115	
63				Cl ₂	+ SO ₂ -	88	-5.4 (25.5)	C ₁₇ H ₁₂ N ₃ O ₂ S ₃	C: 58.39 H: 6.89 N: 7.36 S: 12.64	C: 58.20 H: 7.08 N: 7.32 S: 12.35	3440, 3360, 1662, 1605, 1585, 1510, 1450, 1330, 1290, 1160, 1115, 1095	
64				NH	+ SO ₂ -	74	-23.5 (25.5)	C ₁₈ H ₁₃ N ₃ O ₃ S ₃ ·0.75H ₂ O	C: 50.80 H: 6.88 N: 11.85 S: 11.02	C: 50.98 H: 6.86 N: 11.90 S: 11.46	3380, 2920, 1665, 1605, 1530, 1510, 1405, 1327, 1260, 1153, 1115, 1070	

Table II (continued)

TABLE II (continued)

[18]

Compd. of Ex. No.	R ¹	R ²	R ³	X	R ⁴	Yield%	[α] _D ²⁰ C=1, MeOH(°C)	Molecular formula	Elemental analysis		IR ν _{max} cm ⁻¹
									Calcd.	Found	
65				NH		74	-16.8 (25.5)	C ₂₁ H ₁₉ N ₃ O ₃ S ₂ ·H ₂ O	C: 57.26 H: 7.15 N: 11.40 S: 7.45	C: 57.12 H: 6.93 N: 11.21 S: 7.36	3330, 3000, 1670, 1597, 1525(should), 1507, 1330, 1143, 1124
66				NH		83	-19.9 (25.0)	C ₂₀ H ₁₇ N ₃ O ₃ S ₂ ·H ₂ O ·0.33Cl ₂ Cd ₂	C: 49.92 H: 6.88 N: 11.22 S: 11.00	C: 49.76 H: 6.63 N: 11.10 S: 10.66	3380, 2930, 1665, 1605, 1530, 1510, 1325, 1263, 1155, 1116, 1075, 945
67				NH		86	-14.9 (25.0)	C ₂₂ H ₁₉ N ₃ O ₃ S ₂ ·H ₂ O ·0.5Cl ₂ Cd ₂	C: 55.69 H: 7.04 N: 10.70 S: 7.00	C: 55.48 H: 6.86 N: 10.82 S: 6.66	3330, 2920, 1670, 1600, 1530, 1505, 1446, 1330, 1142, 1115
68				NH		80	-23.8 (25.5)	C ₂₃ H ₁₉ N ₃ O ₃ S ₂ ·H ₂ O	C: 50.17 H: 7.02 N: 12.41 S: 12.17	C: 50.08 H: 6.80 N: 12.41 S: 11.99	3380, 2920, 1665, 1510, 1328, 1262, 1155, 1115
69	Cl ₃			Cl ₂	+SO ₂ -	89	-11.1 (24.0)	C ₂₁ H ₁₈ N ₃ O ₃ S ₂ ·0.5H ₂ O ·0.33Cl ₂ Cd ₂	C: 52.51 H: 6.97 N: 7.58 S: 13.01	C: 52.25 H: 6.80 N: 7.80 S: 12.51	3350, 2920, 1660, 1604, 1525, 1510, 1325, 1286, 1146, 1114
70				Cl ₂	+SO ₂ -	90	-7.0 (23.5)	C ₂₄ H ₁₉ N ₃ O ₃ S ₂ ·0.33H ₂ O	C: 55.72 H: 7.52 N: 7.64 S: 13.12	C: 55.58 H: 7.39 N: 7.57 S: 12.83	3360, 2920, 1660, 1605, 1530, 1510, 1450, 1325, 1290, 1140, 1115

Table 11 (continued)



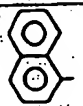
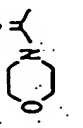
Compd. of Ex. No.	R ¹	R ²	R ³	X	R ⁴	Yield%	[α] _D ²⁰ C=1, MeOH(°C)	Molecular formula	[IR]		
									Elemental analysis		IR ν_{\max} cm ⁻¹
									Calcd.	Found	
71				Cl:		70	-6.6 (24.0)	C ₃₉ H ₃₃ N ₅ O ₇ S ₂ -0.75H ₂ O	C: 59.78 H: 7.27 N: 8.94 S: 8.18	C: 59.81 H: 7.10 N: 8.85 S: 7.89	3340, 2930, 1655, 1628, 1530, 1510, 1447, 1326, 1142, 1126

Table 11 (continued)

Compd. of	[13]	NMR δ
Ex. No. 54		0.74(3H, m), 1.12(6H, m), 1.42(1H, bd), 1.60(3H, bd), 7.51(z), 1.95(2H, m), 2.05(1H, bs), 2.52(2H, m), 2.65(2H, m), 3.00(5H, m), 3.21(5H, m), 3.55(2H, m), 3.78(4H, m), 3.88(1H, m), 4.06~4.25(2H, m), 4.74(1H, m), 5.01(1H, d, J=4Hz), 5.62(1H, t, J=7.5Hz), 6.56(1H, d, J=9.2Hz), 7.18(1H, d, J=1.6Hz), 7.45(1H, m), 7.56~7.76(2H, m), 7.87(1H, dd, J=1.4, 8.2Hz), 7.93(1H, dd, J=1.7, 8Hz), 8.23(1H, d, J=8.4Hz), 8.89(1H, d, J=2Hz), 9.61(1H, d, J=6.8Hz)
55		0.76(3H, m), 0.95~1.53(6H, m), 1.60(1H, bd), 1.80(1H, bs), 2.25(2H, m), 2.42(2H, m), 2.80(1H, m), 3.04(4H, m), 3.19(4H, m), 3.46(7H, m), 3.63(1H, m), 3.75(4H, m), 3.92(1H, dd, J=4.3, 14.3Hz), 3.93(1H, m), 4.51(1H, m), 4.70(1H, m), 5.58(1H, bt), 6.81(1H, d, J=9.5Hz), 7.16(1H, d, J=2.0Hz), 7.46(2H, m), 7.59(2H, m), 7.83(2H, m), 7.89(1H, m), 8.16(1H, d, J=8.2Hz), 8.35(1H, d, J=7.0Hz), 8.61(1H, d, J=1.9Hz)
56		0.60~2.00(13H, m), 2.48(2H, m), 2.58(1H, bs), 2.81(6H, s), 2.68~3.12(4H, m), 3.16~3.63(9H, m), 3.97(2H, m), 4.71(1H, m), 5.27(1H, d, J=5.4Hz), 5.52(1H, bt), 6.57(1H, d, J=9.2Hz), 7.20(1H, d, J=1.8Hz), 7.37(5H, m), 8.90(1H, d, J=1.8Hz), 9.37(1H, d, J=6.8Hz)
57		0.70~1.80(13H, m), 1.34(9H, s), 2.79(6H, s), 2.82~3.53(9H, m), 3.67(1H, m), 3.94(1H, m), 4.67(1H, ddd, J=6Hzx3), 5.73(1H, bt), 6.59(1H, d, J=9.4Hz), 7.27(6H, m), 7.53(1H, d, J=6.6Hz), 8.86(1H, d, J=2)
58		0.75(3H, m), 1.13(5H, m), 1.43(1H, m), 1.60(4H, m), 2.52(4H, m), 2.83(5H, m), 3.22(1H, dd, J=5.15Hz), 3.44(6H, m), 3.87(1H, m), 4.03(1H, m), 4.65(1H, m), 5.33(1H, d, J=5.6Hz), 6.29(1H, t, J=6.3Hz), 6.58(1H, d, J=9.0Hz), 7.16(1H, d, J=1.8Hz), 7.33(6H, m), 7.49(1H, m), 8.20(1H, d, J=8Hz), 8.79(1H, bd), 8.85(1H, d, J=2.0Hz), 9.11(1H, bs), 9.30(1H, d, J=7.0Hz)
59		0.73(3H, bs), 0.92~1.48(6H, m), 1.60(4H, bd), 2.26(2H, m), 2.40(2H, m), 2.73(1H, m), 3.00(4H, m), 3.44(8H, m), 3.85(1H, dd, J=4.2, 14.4Hz), 3.86(1H, m), 4.49(1H, m), 4.63(1H, m), 6.23(1H, bt), 6.80(1H, d, J=9.2Hz), 7.15(1H, d, J=1.6Hz), 7.45(2H, m), 7.58(2H, m), 7.70~7.96(3H, m), 8.16(2H, t, J=9Hz), 8.29(1H, d, J=6.8Hz), 8.58(1H, d, J=1.8Hz), 8.77(1H, m), 9.11(1H, bs)

Table II (continued)

Compd. of Ex. No.	[1 B]	NMR δ
60		0.55~1.70(13H, m), 1.35(9H, s), 2.75(2H, m), 2.85~3.60(8H, m), 3.75(1H, m), 4.55(1H, ddd, J=6.4 Hzx3), 6.17(1H, d, J=9.0Hz), 6.79(1H, t, J=6.7Hz), 7.11(1H, d, J=2.0Hz), 7.26(5H, m), 7.55(1H, dd, J=4.3Hz), 7.65(1H, t, J=7.8Hz), 8.05(1H, dd, J=1.4, 8.2Hz), 8.26(1H, dd, J=1.8, 8.3Hz), 8.41(1H, dd, J=1.4, 7.2Hz), 8.58(1H, d, J=2.0Hz), 9.04(1H, dd, J=1.8, 4.3Hz)
61		0.62~1.75(13H, m), 1.34(9H, s), 2.70~3.54(10H, m), 3.62(1H, m), 3.89(1H, m), 4.61(1H, ddd, J=6.4 Hzx3), 6.40(1H, t, J=6.8Hz), 6.48(1H, d, J=9.2Hz), 7.07(1H, dd, J=3.6, 5Hz), 7.25(6H, m), 7.40(1H, d, J=6.8Hz), 8.70(1H, d, J=2Hz)
62		1.34(9H, m), 0.63~1.78(13H, m), 2.74(1H, dt, J=6.5, 13.5Hz), 2.85~3.52(8H, m), 3.58(1H, dt, J=3.5, 6.6Hz), 4.59(1H, ddd, J=6.5Hzx3), 6.21(1H, t, J=6.4Hz), 6.42(1H, d, J=9.2Hz), 7.19(1H, J=1.7Hz), 7.25(5H, m), 7.41(1H, d, J=6.8Hz), 7.52(3H, m), 7.87(2H, m), 8.66(1H, d, J=2.0Hz)
63		0.60~1.75(13H, m), 1.34(9H, s), 2.63(3H, s), 2.70(1H, dt, J=6.6, 13.6Hz), 2.80~3.46(9H, m), 3.53(1H, m), 3.87(1H, m), 4.47(1H, ddd, J=5.8Hzx3), 5.89(1H, t, J=7Hz), 6.41(1H, d, J=9Hz), 6.92(1H, s), 7.28(5H, m), 7.52(3H, m), 7.63(1H, d, J=5.8Hz), 7.86(2H, dd, J=1.6, 7.7Hz)
64		0.76(3H, m), 1.13(1.43(1H, bd, J=9Hz), 1.60(4H, bd, J=6Hz), 2.02(1H, bs), 2.52(6H, m), 2.72(1H, dd, J=10.16Hz), 2.88(5H, bt, J=7Hz), 3.03(2H, m), 3.26(2H, m), 3.45(8H, m), 3.75(4H, t, J=4.7Hz), 3.92(1H, m), 3.98(1H, m), 4.68(1H, m), 5.17(1H, d, J=5.5Hz), 5.70(1H, bt, J=5Hz), 6.50(1H, d, J=9.6Hz), 7.16(1H, d, J=2.0Hz), 7.34(5H, m), 8.87(1H, d, J=2.0Hz), 9.39(1H, d, J=6.9Hz)
65		0.74(3H, m), 0.9~1.35(5H, m), 1.45(1H, d, J=8Hz), 1.60(4H, m), 1.98(1H, bs), 2.24(2H, m), 2.39(2H, m), 2.55(4H, m), 2.80~3.15(8H, m), 3.25(2H, t, J=7Hz), 3.30~3.68(8H, m), 3.75(4H, m), 3.90(1H, dd, J=4.4, 14.5Hz), 3.91(1H, m), 4.50(1H, m), 4.67(1H, m), 5.67(1H, bt), 6.73(1H, d, J=9.2Hz), 7.15(1H, d, J=1.4Hz), 7.46(2H, m), 7.60(2H, m), 7.76(1H, d, J=3.6Hz), 7.84(1H, dd, J=2.6, 6.8Hz), 7.92(1H, m), 8.16(1H, d, J=8.4Hz), 8.34(1H, d, J=7.2Hz), 8.60(1H, d, J=2Hz)

Table II (continued)

Compd. of	[13C]	NMR δ
Ex. No. 66		0.55~1.74(13H, m), 2.02(2H, m), 2.18(1H, bs), 2.50(6H, m), 2.66~2.94(3H, m), 2.94~3.30(5H, m), 3.45(8H, m), 3.73(5H, m), 3.90(1H, m), 4.00(1H, m), 4.67(1H, m), 5.20(1H, bd), 5.78(1H, bt), 6.53(1H, d, J=9.4Hz), 7.16(1H, d, J=1.9Hz), 7.35(5H, m), 8.87(1H, d, J=2.0Hz), 9.41(1H, d, J=6.6Hz)
67		0.62~1.75(13H, m), 2.04(2H, m), 2.20(2H, m), 2.39(2H, m), 2.51(6H, m), 2.90(3H, m), 3.10(4H, m), 3.43(6H, m), 3.57(2H, m), 3.73(4H, m), 3.90(2H, m), 4.50(1H, m), 4.69(1H, m), 5.76(1H, bt), 6.77(1H, d, J=9.1Hz), 7.16(1H, d, J=1.4Hz), 7.44(2H, m), 7.60(2H, m), 7.76(1H, d, J=3.2Hz), 7.85(1H, m), 7.90(1H, m), 8.16(1H, d, J=8.1Hz), 8.37(1H, d, J=6.8Hz), 8.60(1H, d, J=2.1Hz)
68		0.73(3H, m), 1.15(5H, m), 1.43(1H, bd, J=8.1Hz), 1.61(4H, bd, J=6.1Hz), 2.31(6H, s), 2.50(2H, m), 2.74(1H, dd, J=10.14Hz), 2.83(5H, m), 3.04(2H, m), 3.26(2H, m), 3.45(7H, m), 3.90(1H, m), 4.02(1H, dd, J=2.8, 10.4Hz), 4.69(1H, m), 5.23(1H, bs), 6.51(1H, d, J=9.1Hz), 7.17(1H, d, J=1.6Hz), 7.35(5H, m), 7.35(5H, m), 8.87(1H, d, J=2.1Hz), 9.32(1H, d, J=7.1Hz)
69		0.70~1.80(13H, m), 1.35(9H, s), 2.96(3H, s), 2.75(1H, bs), 2.87(~3.50(9H, m), 3.65(1H, m), 3.45(1H, m), 4.63(1H, ddd, J=5.8Hz), 5.78(1H, t, J=6.6Hz), 6.50(1H, d, J=9.2Hz), 7.28(6H, m), 7.60(1H, d, J=2.1Hz), 8.77(1H, d, J=2.1Hz)
70		0.70~1.88(17H, m), 0.95(3H, t, J=7.2Hz), 2.87~3.52(12H, m), 3.63(1H, m), 3.94(1H, m), 4.63(1H, ddd, J=6.2Hzx3), 5.68(1H, t, J=6.4Hz), 6.45(1H, d, J=9.1Hz), 7.25(6H, m), 7.54(1H, d, J=6.4Hz), 8.76(1H, d, J=2.1Hz)
71		0.64~1.88(17H, m), 0.94(3H, t, J=7.2Hz), 2.28(1H, dd, J=6.4, 16.6Hz), 2.60~3.80(19H, m), 4.04(1H, m), 4.70(1H, ddd, J=4.7Hzx3), 5.57(1H, t, J=6.8Hz)

Renin inhibition potency of the compounds (I) of the invention was determined in vitro and in vivo according to the procedure described in the following Experiments.

Experiment 1 Potency in vitro

Commercially available lyophilized human plasma (Ortho, Bi-Level Plasma Renin Control) was renatured by dissolving in water. Angiotensinogen was allowed to react with intrinsic renin contained in the renatured plasma to generate angiotensin I (AI), which was quantitatively measured with radioimmunoassay (RIA). Thus, potency of the plasma renin was determined on the basis of the AI production. For this purpose, Renin RIA kit (RENIN[®] RIABEAD[®]) manufactured by Dinabott was used. All of the reagents necessary for the measurement of the AI production were available from the attachment of the kit, and the measurement was conducted according to the manufacturer's direction.

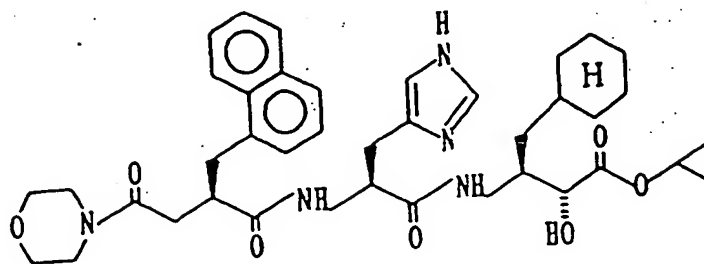
To the plasma (0.2ml) were added all of the reagents, and the mixture was combined with either of sample solutions (0.002ml) of various concentrations which had been prepared by dissolving a test compound in different amount of ethanol. Ethanol (0.002ml) containing no test compound was used as a control solution. The amount of AI produced was measured after 60 minutes incubation. Renin inhibition potency of test compound was determined by comparing the amount of AI produced by a sample solution with that produced by a control solution. The concentrations of the test compounds which inhibit renin activity by 50% (IC_{50}) are summarized in Table 11.

Table 11 Renin Inhibition in vitro

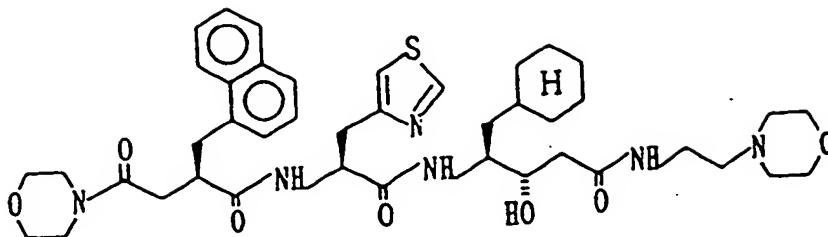
Test Compound IC ₅₀ (Example No.)	Test Compound IC ₅₀ (Example No.)	Test Compound IC ₅₀ (Example No.)
1 6.09	22 39.2	42 13
2 5.87	23 2.07	43 0.51
3 4.44	24 1.56	44 1.53
4 3.21	25 3.17	45 0.31
5 29.0	26 1.32	46 3.16
6 4.22	27 1.78	47 5.90
8 6.17	28 0.52	48 1.98
9 12.0	29 3.31	49 2.34
10 10.9	30 1.07	50 14.8
11 9.1	31 11.6	51 4.51
12 4.56	32 6.72	52 1.69
13 53.9	33 4.65	53 0.36
14 9.3	34 9.53	55 0.60
15 12.6	35 0.63	56 0.70
16 71.3	36 4.98	57 0.80
17 259	37 14.5	58 0.19
18 22.8	38 39.2	59 0.41
19 3.75	39 7.52	62 1.24
20 7.36	40 18.1	64 0.70
21 2.73	41 4.98	69 0.53
		(1)(KRI-1314) 21.3
		(2)(ES-6864) 3.75

IC₅₀ : nM

(1) KRI-1314



(2) ES-6864

Experiment 2 Potency in vivo

Crab-eating monkeys (*Cynomolgus* monkeys) (2.8-5.0 kg) were fed on low sodium diet (Na 7.15mg/100g feed) for six days, during which the monkeys intramuscularly received furosemide (2mg/kg body weight) every other day from the second day of the experiment, in order to make the monkeys hyperrenin condition.

After seven days of low sodium feeding, the monkeys were restrained on a monkey chair. Compounds to be tested are dissolved in 0.1M citric acid/physiological saline or suspended in water with addition of β -cyclodextrin, and orally administered to the monkeys using a stomach probe (15mg/kg body weight). Two milliliters of blood was collected from the femoral vein before administration of the compounds and 0.5, 1.5, 2.5 and 4 hours after the administration. For the blood collection, an injection syringe containing 30 μ l of 6% aqueous EDTA-2Na solution was used. The collected blood was transferred into a test tube and centrifuged (3000 rpm, 10 minutes) at 4°C, and the resultant supernatant was used to determine the renin content. Plasma renin activity (AI(ng)/ml/h value) was measured using a Radioimmunoassay kit commercially available from Dinabott Co. in the same manner as in the foregoing *in vitro* test. Renin inhibition potencies of the compounds tested, which were expressed as a percentage of renin activity relative to the activity before the administration, are listed in Table 12.

Table 12

Compound	Max	Mean	4h	6h	8h	24h
Example No.						
1	33	22	22			
2	49	46	49			
8	60	52	60			
21	55	37	55			
24	99	90	77	56	42	28
26	83	71	69	99		
27	81	65	81	73	64	28
28	97	74	97	89	83	53
33	95	85	68	82		
35	39	30	39	23	24	14
39	46	28	12			
40	44	30	44			
41	95	89	87	76	54	18
43	98	86	95	83	70	11
44	99	97	91	81	71	21
47	59	47	55	6	18	34
48	98	88	88	63	33	0
49	92	84	78	72	51	12
50	93	59	58	42	29	0
51	80	48	35	0	0	0
53	96	85	94		73	
56	93	72	82		85	
57	100	97	89		80	
58	83	71	82		67	

1) Administration rate of compound No. 1 is 30mg/kg.

2) Furosemide was not administered in case of Nos. 2 and 8.

The compounds of the invention which are not listed in Table 12 showed similar inhibition potencies.

Vasodepressor activity of the compounds of the invention was also measured with direct technique using a conscious monkey, where a monkey was administered a compound of the invention orally or intravenously (a solution in Tween 20). The test results are shown in Table 13.

Table 13

Compound Example No.	Administration route	Dose (mg/kg)	Maximum reduced BP (-ΔmmHg)
43	p.o.	100	35
		30	10
		10	5
43	i.v.	3	-
		1	20
		0.3	5
44	i.v.	3	20
		1	8
		0.3	5

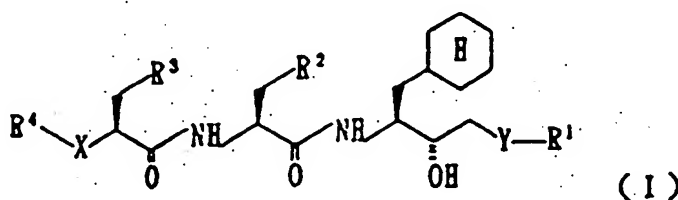
The above test results show that the compounds of the present invention have renin inhibition potency both *in vitro* and *in vivo*.

The compounds of the invention are thus useful for the treatment of hypertension due to the renin inhibition when orally administered. However, other administration routes may be also effective.

As discussed previously, the compounds of the invention can be formulated into a pharmaceutical composition together with suitable carriers or excipients. When the compounds of the invention are used as a hypotensive agent, suitable dosage is 0.01-50mg/kg/day in one to three divided doses, preferably 0.05-10mg/kg/day, when orally administered, and 1-5000μg/kg/day, preferably 5-500μg/kg/day, when parenterally administered.

Claims

1. A dipeptide derivative of formula (I):



wherein:

R¹ is C₁-C₁₂ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₁₀ cycloalkyl, aryl, or heterocyclic radical;
 R² is carbamoyl, aryl, 5- or 6-membered heterocyclic radical, C₁-C₁₂ alkyl-S-, C₁-C₁₂ alkyl-S-CH₂-, or C₃-C₁₀ cycloalkyl-S-;

R³ is aryl of 5- or 6-membered heterocyclic radical;

R⁴ is R⁴-SO₂ or R⁴-CO;

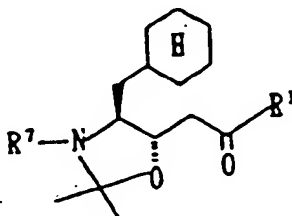
R⁴ is aryl, C₁-C₁₂ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl; C₃-C₁₀ cycloalkyl, or heterocyclic radical;

X is CH₂, NH, O, or S; and

Y is CO or NHSO₂ wherein R¹, R², R³ and R⁴ each may be substituted with one to three substituents selected independently from a group consisting of hydroxy; halogen; trifluoromethyl; -CN; heterocyclic radical; C₁-C₆ alkyl; C₃-C₁₀ cycloalkyl; -O-C₁-C₆ alkyl; -S-C₁-C₆ alkyl; -SO-C₁-C₆ alkyl; -SO₂-C₁-C₆ alkyl; C₁-C₆ alkylendioxy; -CO-O-C₁-C₆ alkyl; -NHCO-C₁-C₆ alkyl; -NHSO₂-C₁-C₆ alkyl; -NR⁵R⁶; -O-CO-NR⁵R⁶; -CO-NR⁵R⁶; -O-C₁-C₆ alkyl NR⁵R⁶; R⁵ and R⁶ are independently hydrogen, formyl or C₁-C₆ alkyl, or R⁵ and

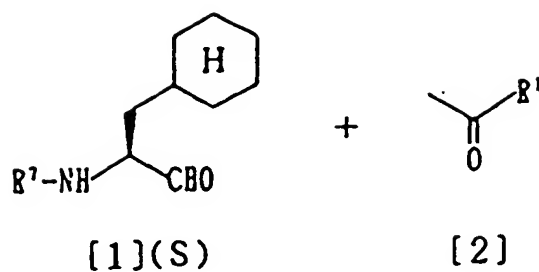
R^6 , when taken together with the nitrogen to which they are attached, form a cyclic amino group; or an acid addition salt thereof.

2. A compound as claimed in Claim 1 wherein R^2 is optionally substituted 5- or 6-membered heterocyclic group; R^3 is optionally substituted aryl; R^4 is morpholinosulfonyl; and x is NH.
3. A compound for the manufacture of the derivative of formula (I) of Claim 1, said compound having the formula:

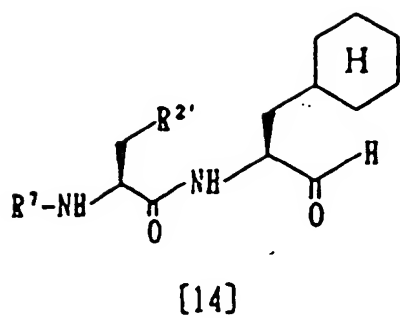


wherein, R^1 is as defined in Claim 1, and R^7 is hydrogen or an amino protecting group.

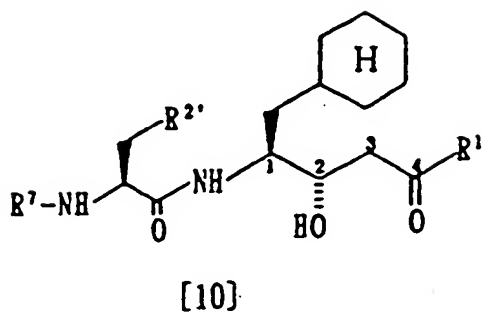
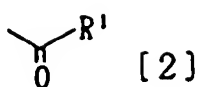
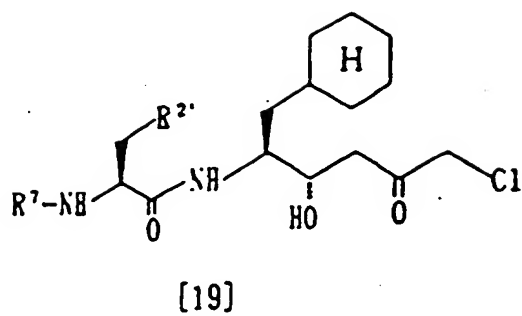
4. A process for the preparation of a compound as defined in Claim 1 of Claim 2 wherein Y is CO comprising at least the final step of the following reaction scheme:

Step 1

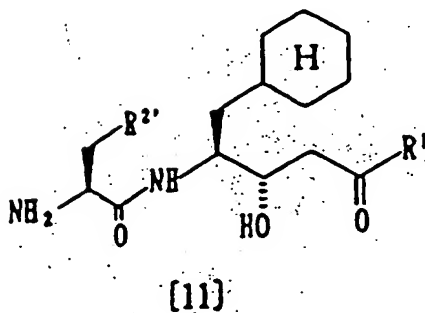
Step 2b



Step 2c

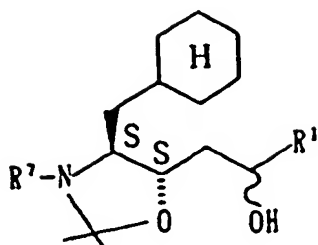


Step 3

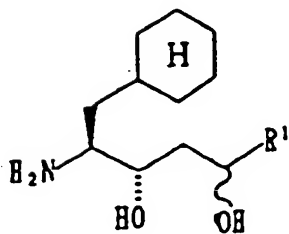


Step 2a

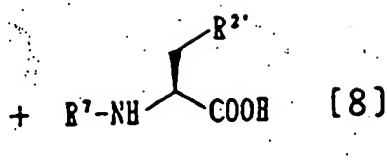
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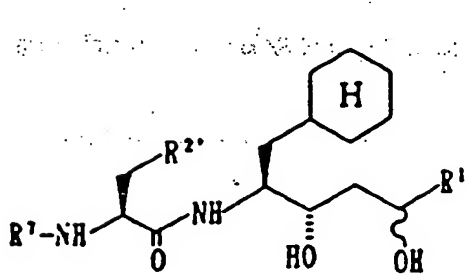
[6]



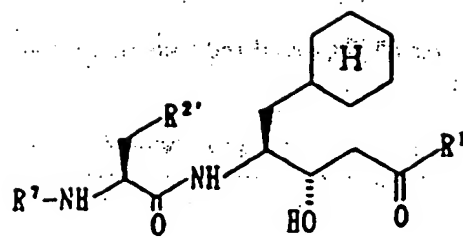
[7]



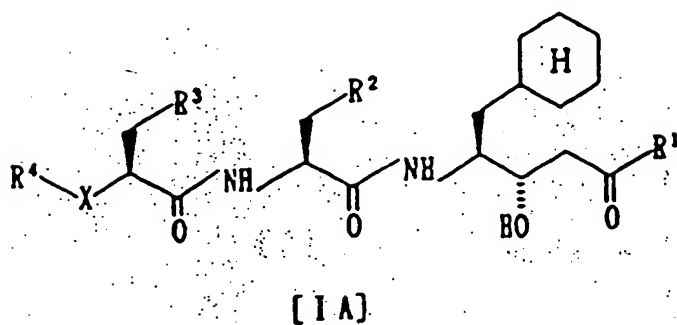
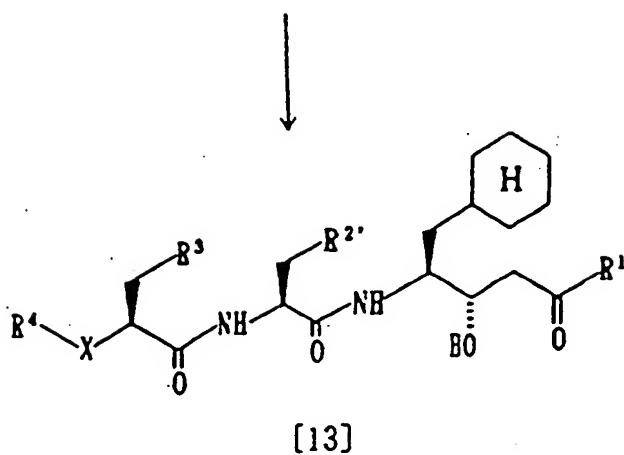
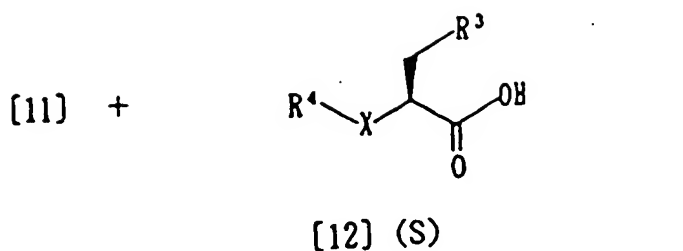
[8]



[9]



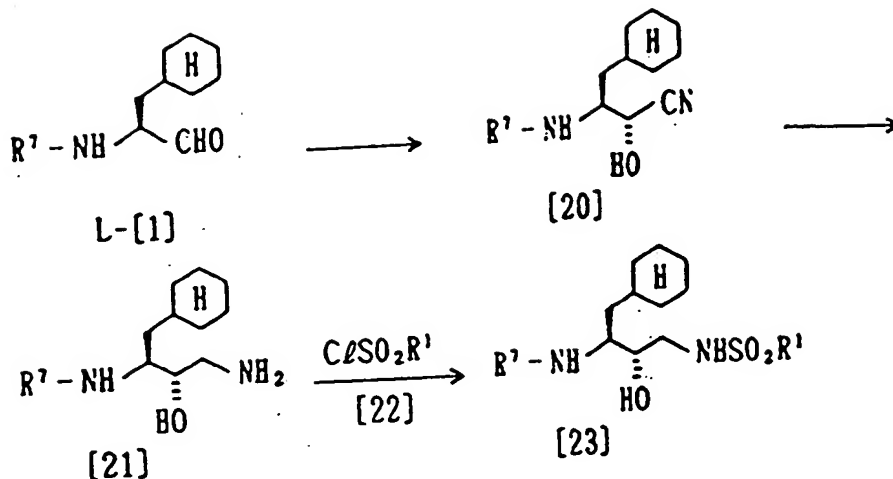
[10]

Step 4

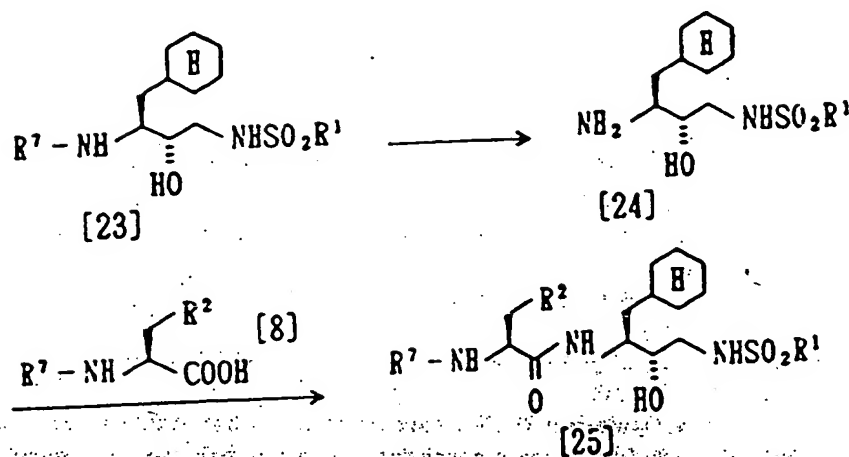
wherein R¹, R², R³, R⁴ and X are as defined in Claim 1, R^{2'} is protected R² and R⁷ is an amino protecting group.

5. A process for preparing a compound as defined in Claim 1 or Claim 2 wherein Y is NHSO₂ comprising at least the final step of the following reaction scheme:

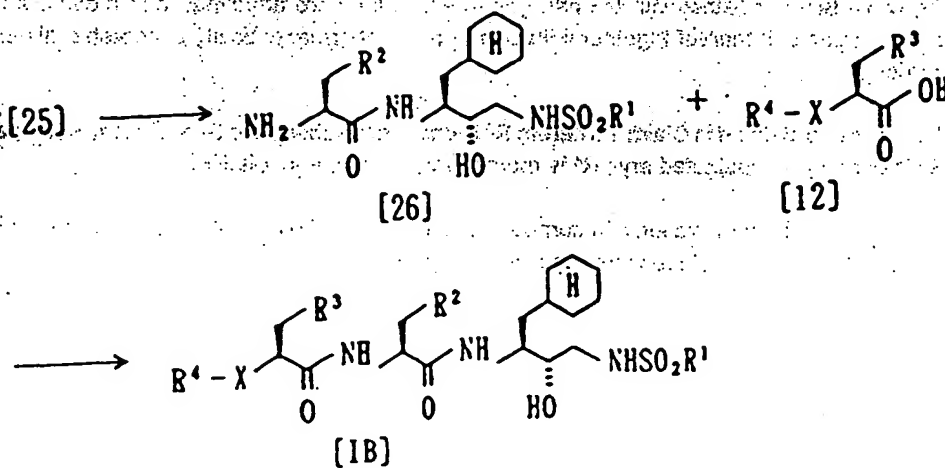
Step 1



Step 2



Step 3



wherein R¹, R², R³, R⁴ and X are as defined in Claim 1 and R⁷ is an amino protecting group.

6. A pharmaceutical preparation for use in the treatment of hypertension comprising a pharmaceutically effective amount of at least one compound as defined in Claim 1 or Claim 2 together with one or more pharmaceutically acceptable carriers, diluents or excipients.

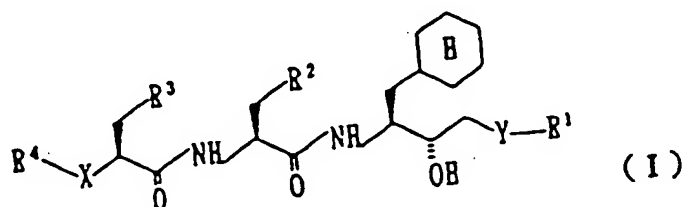
7. The use of a compound as defined in Claim 1 or Claim 2 in the manufacture of a medicament for use in the treatment of hypertension.

8. A process for carrying out a stereo selective aldol condensation between an aldehyde and a ketone wherein the reaction is carried out in the presence of a metal amide and a crown ether in an organic solvent and at a temperature in the range -10 to about -100°C.

9. A process as claimed in Claim 8 wherein the amide is sodium bis-trimethylsilylamide ($\text{NaN}(\text{TMS})_2$), the crown ether is 15-crown-5 and the temperature is about -78°C.

Claims for the following Contracting State : ES

1. A process for the production of a pharmaceutical preparation for the treatment of hypertension comprising the step of admixing a pharmaceutically effective amount of at least one compound of the formula



wherein:

R^1 is C_1 - C_{12} alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_{10} cycloalkyl, aryl, or heterocyclic radical;

R^2 is carbamoyl, aryl, 5- or 6-membered heterocyclic radical, C_1 - C_{12} alkyl-S-, C_1 - C_{12} alkyl-S- CH_2 -, or C_3 - C_{10} cycloalkyl-S-;

R^3 is aryl of 5- or 6-membered heterocyclic radical;

R^4 is R^4 - SO_2 or R^4 -CO;

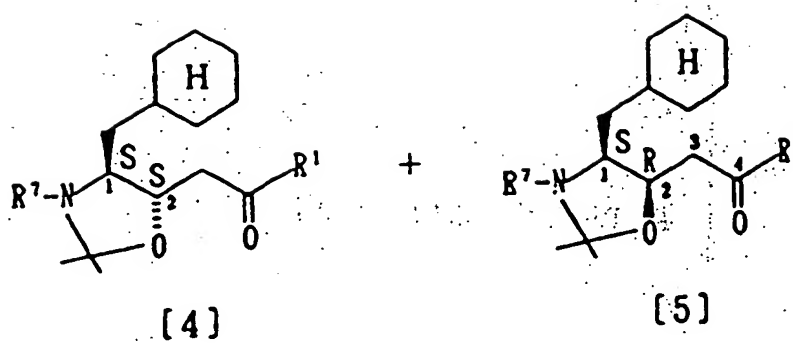
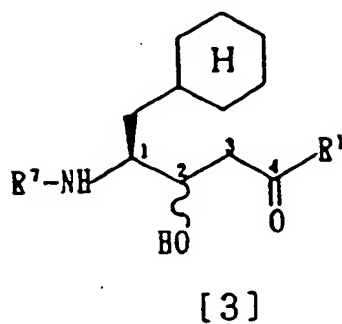
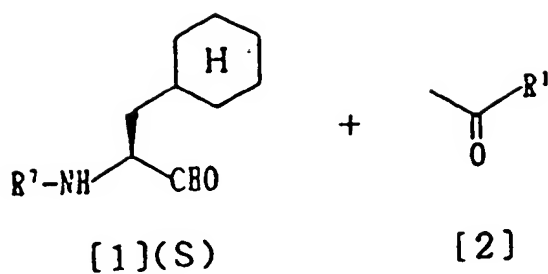
R^4 is aryl, C_1 - C_{12} alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_{10} cycloalkyl, or heterocyclic radical;

X is CH_2 , NH, O, or S; and

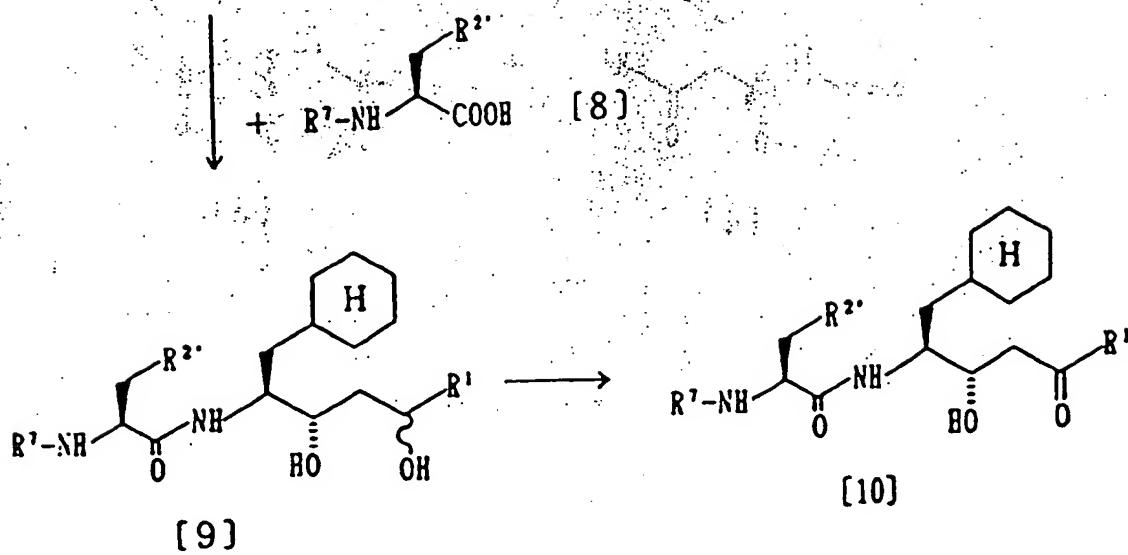
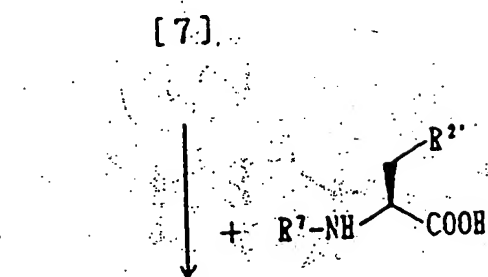
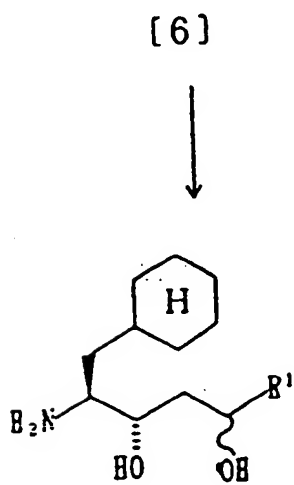
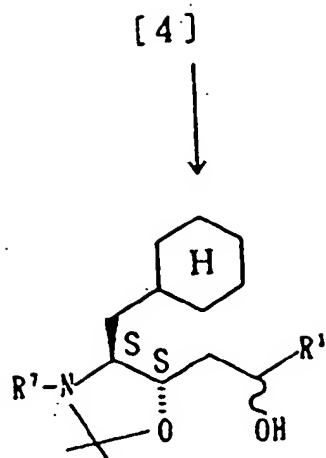
Y is CO or NHSO_2 wherein R^1 , R^2 , R^3 and R^4 each may be substituted with one to three substituents selected independently from a group consisting of hydroxy; halogen; trifluoromethyl; -CN; heterocyclic radical; C_1 - C_8 alkyl; C_3 - C_{10} cycloalkyl; -O- C_1 - C_8 alkyl; -S- C_1 - C_8 alkyl; - SO_2 - C_1 - C_8 alkyl; C_1 - C_8 alkylenedioxy; -CO-O- C_1 - C_8 alkyl; -NHCO- C_1 - C_8 alkyl; - NHSO_2 - C_1 - C_8 alkyl; - NR^5R^6 ; -O-CO- NR^5R^6 ; -CO- NR^5R^6 ; -O- C_1 - C_8 alkyl NR^5R^6 ; R^5 and R^6 are independently hydrogen, formyl or C_1 - C_8 alkyl, or R^5 and R^6 , when taken together with the nitrogen to which they are attached, form a cyclic amino group; or an acid addition salt thereof together with one or more pharmaceutically acceptable diluents, excipients or carriers.

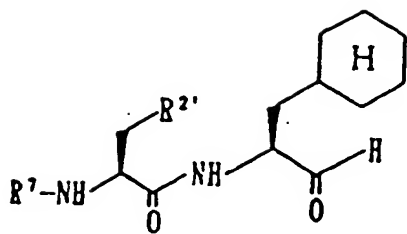
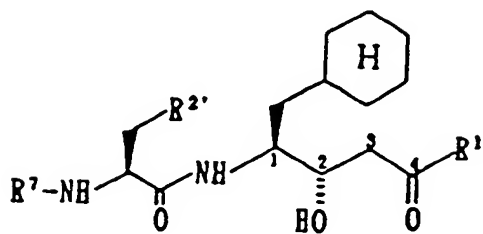
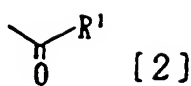
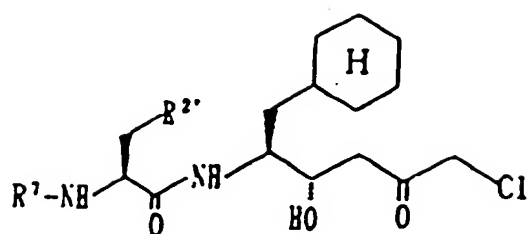
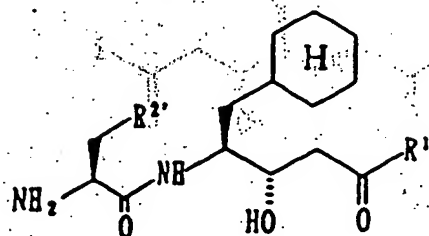
2. A process as claimed in Claim 1 wherein R^2 is optionally substituted 5- or 6-membered heterocyclic group; R^3 is optionally substituted aryl; R^4 is morpholinosulfonyl; and x is NH.

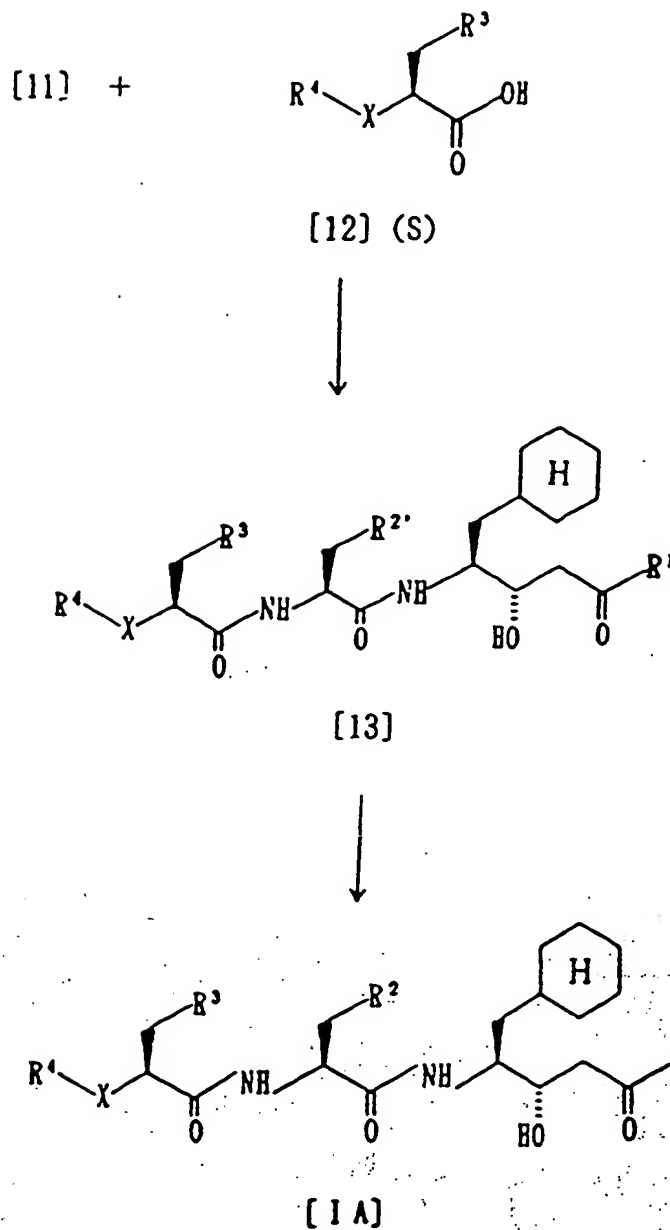
3. A process for the preparation of a compound as defined in Claim 1 of Claim 2 wherein Y is CO comprising at least the final step of the following reaction scheme:

Step 1

Step 2a



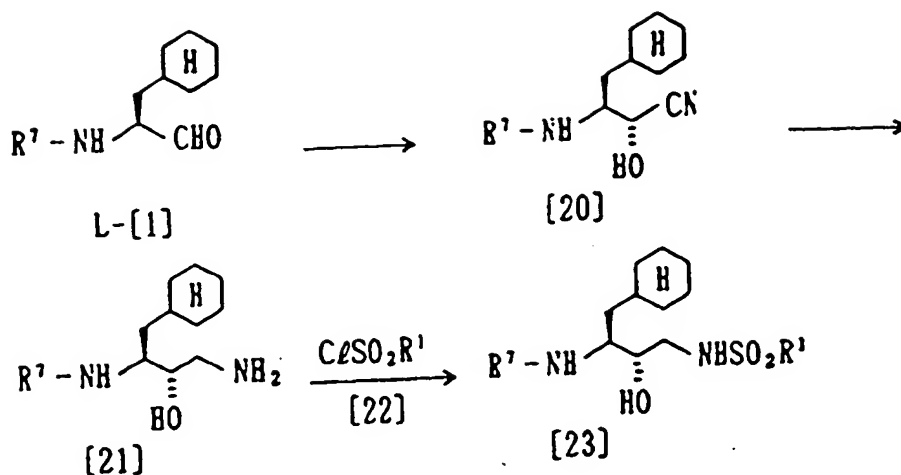
Step 2bStep 2cStep 3

Step 4

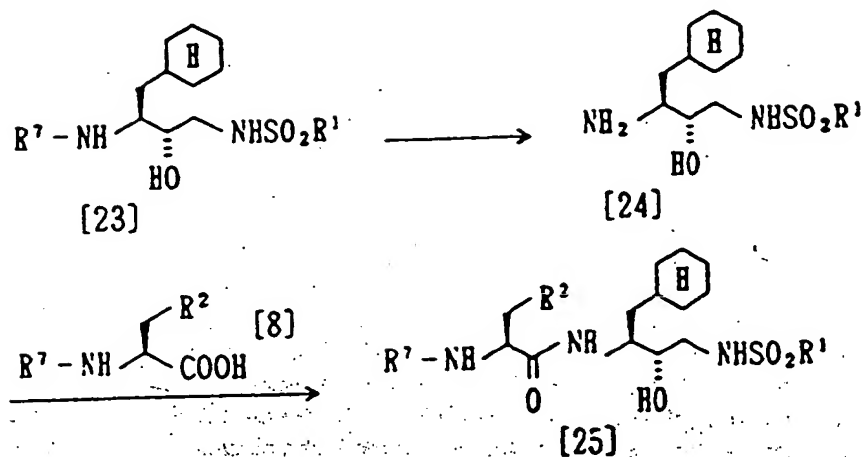
wherein R^1 , R^2 , R^3 , R^4 and X are as defined in Claim 1, $\text{R}^{2'}$ is protected R^2 and R^7 is an amino protecting group.

4. A process for preparing a compound as defined in Claim 1 or Claim 2 wherein Y is NHSO_2 comprising at least the final step of the following reaction scheme:

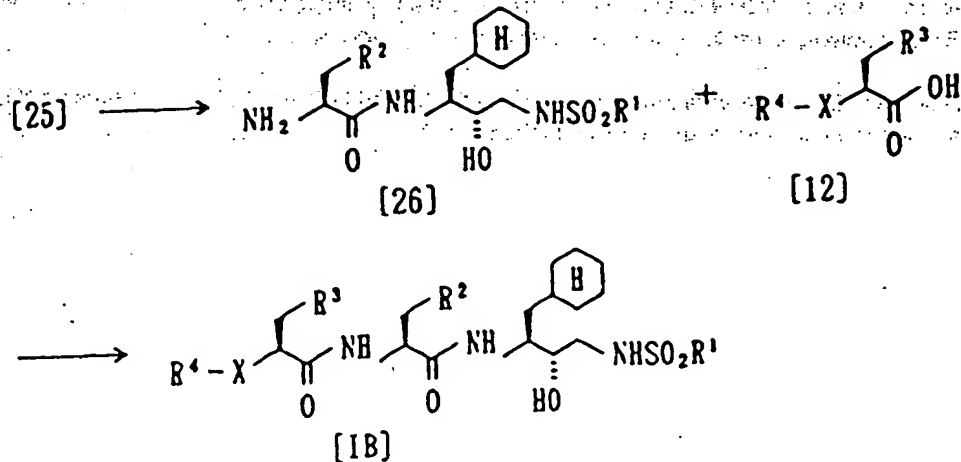
Step 1



Step 2



Step 3



wherein R¹, R², R³, R⁴ and X are as defined in Claim 1 and R⁷ is an amino protecting group.

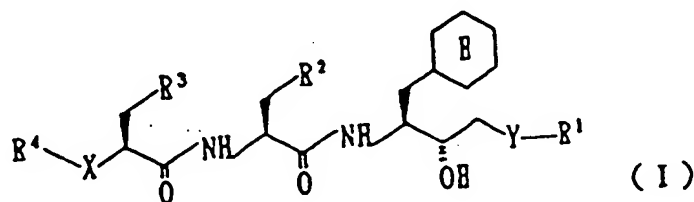
5. The use of a compound as defined in Claim 1 or Claim 2 in the manufacture of a medicament for use in

the treatment of hypertension.

6. A process for carrying out a stereo selective aldol condensation between an aldehyde and a ketone wherein the reaction is carried out in the presence of a metal amide and a crown ether in an organic solvent and at a temperature in the range -10 to about -100°C.
7. A process as claimed in Claim 8 wherein the amide is sodium bis-trimethylsilylamide ($\text{NaN}(\text{TMS})_2$), the crown ether is 15-crown-5 and the temperature is about -78°C.

Claims for the following Contracting States: GR

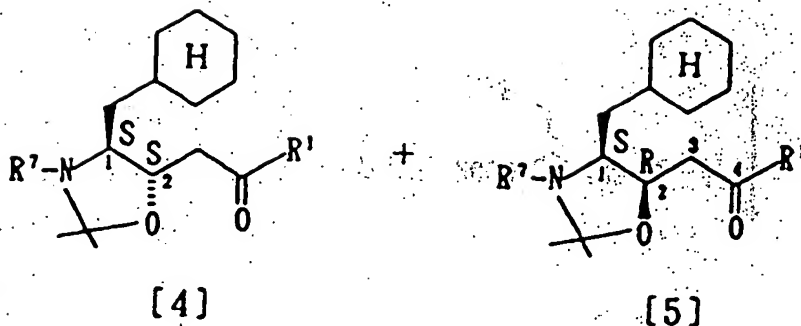
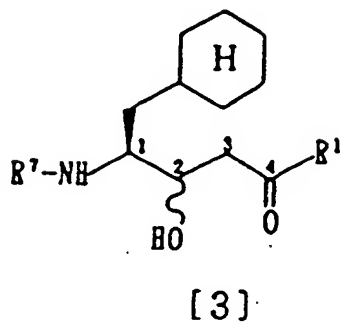
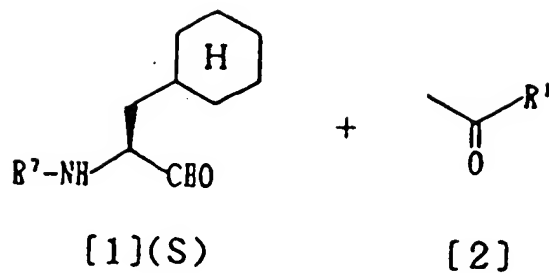
1. A process for the production of a pharmaceutical preparation for the treatment of hypertension comprising the step of admixing a pharmaceutically effective amount of at least one compound of the formula

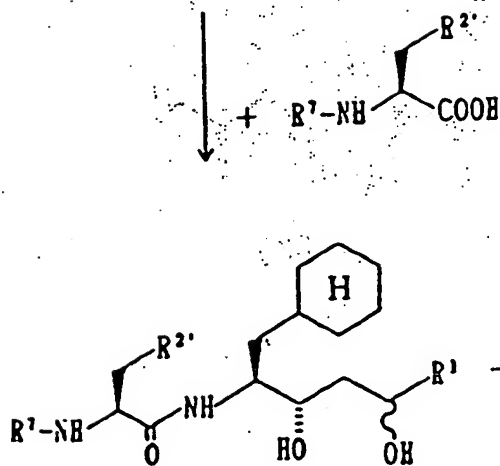
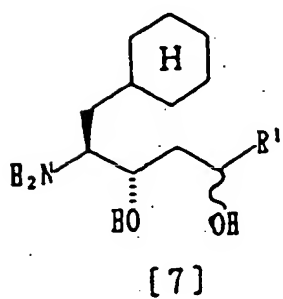
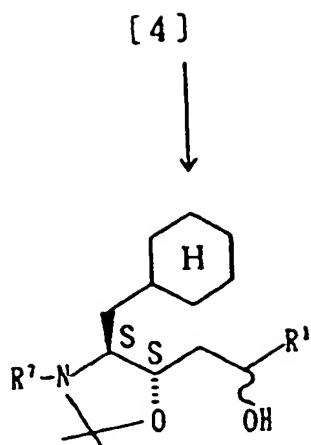


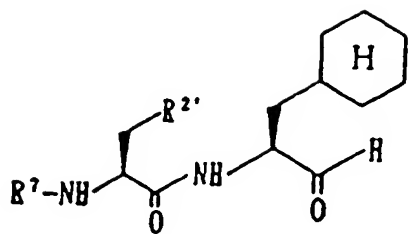
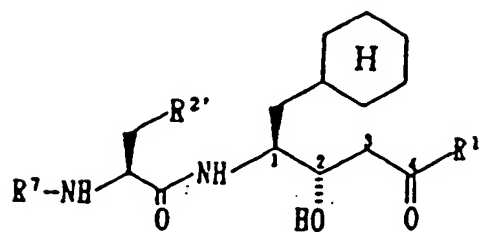
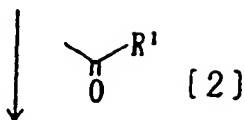
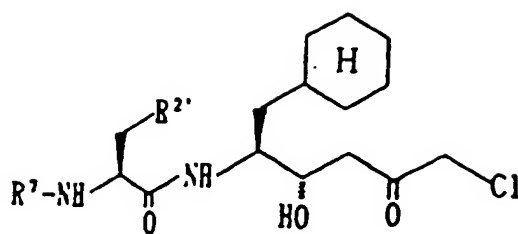
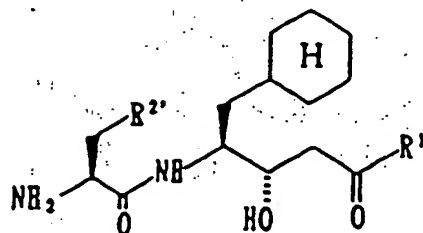
wherein:

- R^1 is C_1 - C_{12} alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_{10} cycloalkyl, aryl, or heterocyclic radical;
- R^2 is carbamoyl, aryl, 5- or 6-membered heterocyclic radical, C_1 - C_{12} alkyl-S-, C_1 - C_{12} alkyl-S- CH_2 -, or C_3 - C_{10} cycloalkyl-S-;
- R^3 is aryl or 5- or 6-membered heterocyclic radical;
- R^4 is R^4 - SO_2 or R^4 -CO;
- R^4 is aryl, C_1 - C_{12} alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_{10} cycloalkyl, or heterocyclic radical;
- X is CH_2 , NH, O, or S; and
- Y is CO or NHSO_2 wherein R^1 , R^2 , R^3 and R^4 each may be substituted with one to three substituents selected independently from a group consisting of hydroxy; halogen; trifluoromethyl; -CN; heterocyclic radical; C_1 - C_6 alkyl; C_3 - C_{10} cycloalkyl; -O- C_1 - C_6 alkyl; -S- C_1 - C_6 alkyl; -SO- C_1 - C_6 alkyl; - SO_2 - C_1 - C_6 alkyl; C_1 - C_6 alkylenedioxy; -CO-O- C_1 - C_6 alkyl; -NHCO- C_1 - C_6 alkyl; - NHSO_2 - C_1 - C_6 alkyl; - NR^5R^6 ; -O-CO- NR^5R^6 ; -CO- NR^5R^6 ; -O- C_1 - C_6 alkyl NR^5R^6 ; R^5 and R^6 are independently hydrogen, formyl or C_1 - C_6 alkyl, or R^5 and R^6 , when taken together with the nitrogen to which they are attached, form a cyclic amino group; or an acid addition salt thereof together with one or more pharmaceutically acceptable diluents, excipients or carriers.

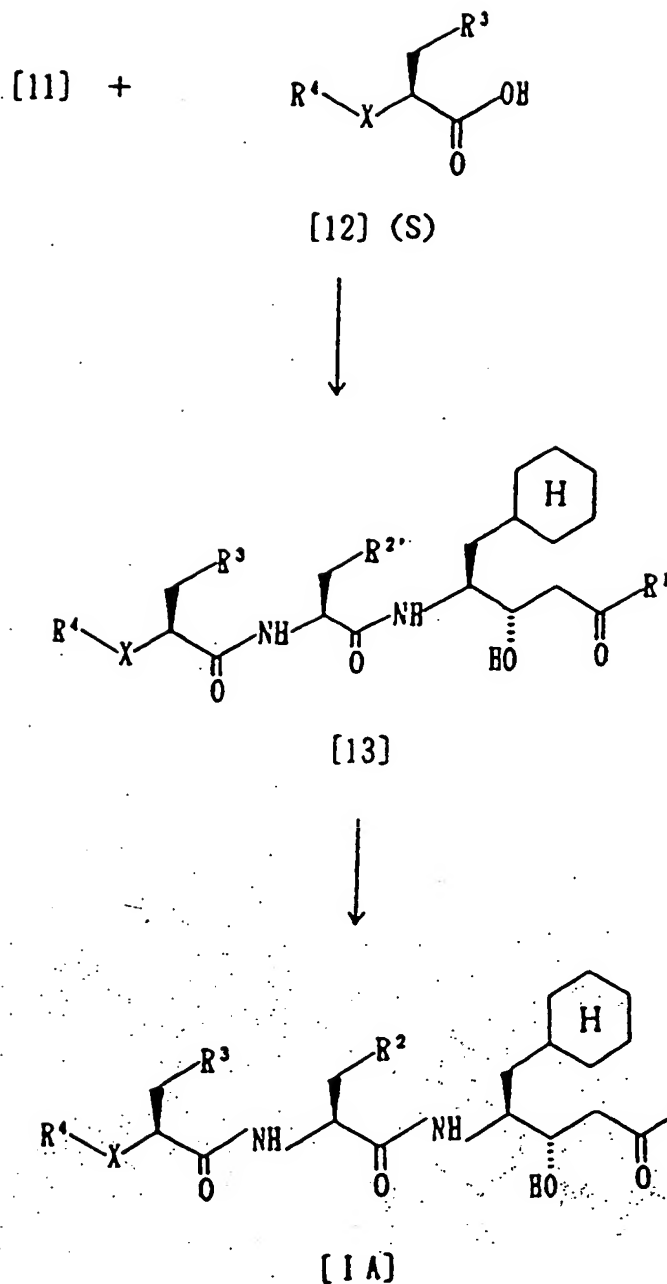
2. A process as claimed in Claim 1 wherein R^2 is optionally substituted 5- or 6-membered heterocyclic group; R^3 is optionally substituted aryl; R^4 is morpholinosulfonyl; and x is NH.
3. A process for the preparation of a compound as defined in Claim 1 of Claim 2 wherein Y is CO comprising at least the final step of the following reaction scheme:

Step 1

Step 2a

Step 2bStep 2cStep 3

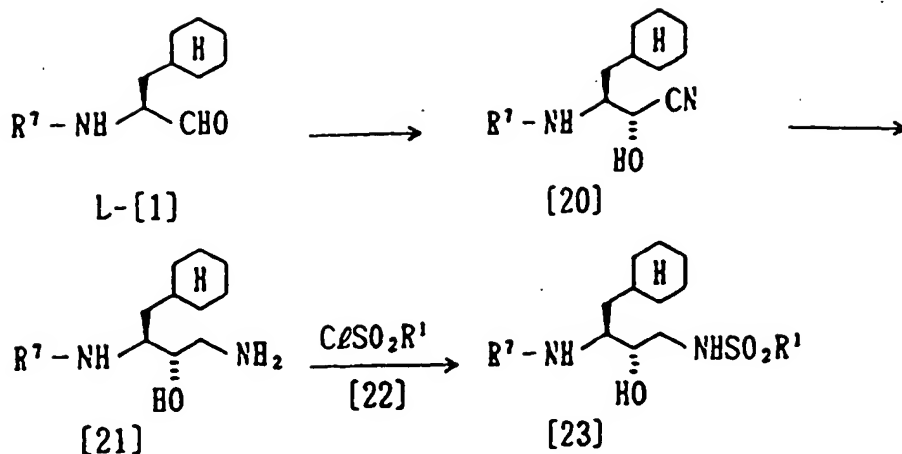
[11]

Step 4

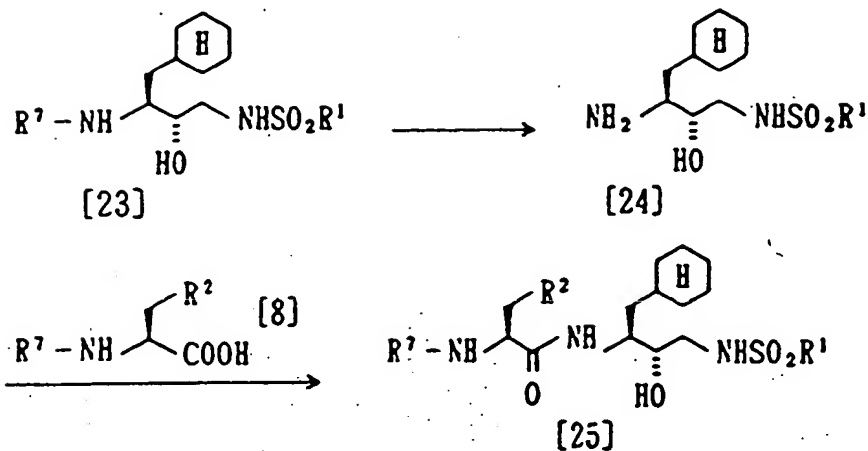
wherein R^1 , R^2 , R^3 , R^4 and X are as defined in Claim 1, R^2 is protected R^2 and R^7 is an amino protecting group.

4. A process for preparing a compound as defined in Claim 1 or Claim 2 wherein Y is NHSO_2 comprising at least the final step of the following reaction scheme:

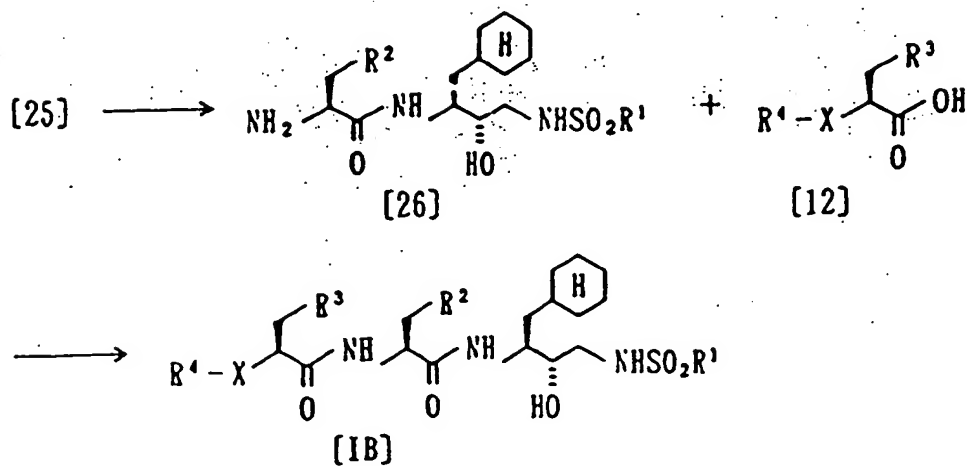
Step 1



Step 2

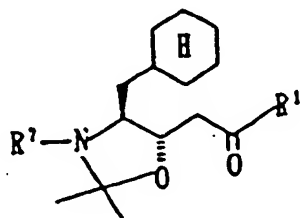


Step 3



wherein R¹, R², R³, R⁴ and X are as defined in Claim 1 and R⁷ is an amino protecting group.

5. The use of a compound as defined in Claim 1 or Claim 2 in the manufacture of a medicament for use in the treatment of hypertension.
6. A process for carrying out a stereo selective aldol condensation between an aldehyde and a ketone wherein the reaction is carried out in the presence of a metal amide and a crown ether in an organic solvent and at a temperature in the range -10 to about -100°C.
7. A process as claimed in Claim 8 wherein the amide is sodium bis-trimethylsilylamide ($\text{NaN}(\text{TMS})_2$), the crown ether is 15-crown-5 and the temperature is about -78°C.
8. A compound for the manufacture of the derivative of formula (I) of Claim 1, said compound having the formula:



wherein, R¹ is as defined in Claim 1, and R² is hydrogen or an amino protecting group.